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OM protein - protein search, using sw model

Run on: March 1, 2004, 10:03:28 ; Search time 16.5 Seconds
(without alignments)
137.669 Million cell updates/sec

Title: SEQ1-4SUBS

Perfect score: 1 ANSFLXXLRGSLXRCIXX.....XXAKIFEDVDTLAFWSKH 44

Sequence: BLOSUM62

Scoring table: Gapop 10.0, Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: Issued_Patents_AA.*

1: /cgn2_6/ptodata/2/1aa/5A.COMB.pep.*
2: /cgn2_6/ptodata/2/1aa/5B.COMB.pep.*
3: /cgn2_6/ptodata/2/1aa/6A.COMB.pep.*
4: /cgn2_6/ptodata/2/1aa/6B.COMB.pep.*
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6: /cgn2_6/ptodata/2/1aa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARY

Result No.	Score	Query Match Length	ID	Description
1	179	90.9	44	US-08-955-636-24
2	179	90.9	419	US-10-182-263-6
3	176	89.3	44	US-08-955-636-35
4	174	88.3	419	US-10-182-263-3
5	174	88.3	419	US-10-182-263-4
6	174	88.3	419	US-10-182-263-5
7	173	87.8	44	US-08-955-636-20
8	170	86.3	44	US-08-955-636-21
9	168	85.3	44	US-08-955-636-19
10	168	85.3	44	US-08-955-636-22
11	160	81.2	44	US-08-955-636-1
12	160	81.2	44	US-08-955-636-25
13	160	81.2	45	US-08-955-636-2
14	160	81.2	419	US-08-955-411-1
15	160	81.2	419	US-08-955-411-1
16	160	81.2	419	US-09-667-570A-3
17	160	81.2	419	US-10-182-263-1
18	160	81.2	419	PCT-US92-10242-1
19	160	81.2	460	US-08-756-506-4
20	160	81.2	460	US-08-756-506-4
21	160	81.2	460	5270178-13
22	160	81.2	460	5270178-14
23	160	81.2	460	5270178-15
24	160	81.2	461	5270178-16
25	160	81.2	461	US-10-182-263-2
26	160	81.2	461	5225537-2
27	160	81.2	461	5270178-17

28	160	81.2	461	5270178-18	Patent No. 5270178
29	160	81.2	461	5460953-3	Patent No. 5460953
30	147	74.6	42	US-08-745-254A-2	Sequence 2, Appl.1
31	147	74.6	461	5270178-2	Patent No. 5270178
32	143	72.6	41	US-08-229-280-5	Sequence 5, Appl.1
33	138	70.1	42	US-09-383-667-8	Sequence 8, Appl.1
34	129	65.5	409	US-09-065-872-2	Sequence 2, Appl.1
35	129	65.5	409	US-09-667-570A-2	Sequence 2, Appl.1
36	129	65.5	410	US-09-065-872-1	Sequence 1, Appl.1
37	129	65.5	410	US-09-667-570A-1	Sequence 1, Appl.1
38	117	59.4	44	US-08-955-636-23	Sequence 23, Appl.1
39	116	58.9	44	US-08-955-636-2	Sequence 2, Appl.1
40	114	57.9	139	US-08-330-978-2	Sequence 2, Appl.1
41	114	57.9	139	US-08-474-042-2	Sequence 2, Appl.1
42	114	57.9	139	US-08-484-558-2	Sequence 2, Appl.1
43	114	57.9	139	US-08-774-592-2	Sequence 2, Appl.1
44	114	57.9	437	US-08-487-037-2	Sequence 2, Appl.1
45	114	57.9	437	US-08-487-037-3	Sequence 3, Appl.1

ALIGNMENTS

```
RESULT 1
US-08-955-636-24
Sequence 24, Application US/08955636A
Patent No. 6017892
GENERAL INFORMATION:
APPLICANT: Nelsentuen, Gary
TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
FILE REFERENCE: 09531/002001
CURRENT APPLICATION NUMBER: US/08/955,636A
CURRENT FILING DATE: 1997-10-23
NUMBER OF SEQ ID NOS: 35
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 24
LENGTH: 44
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: MOD RES
LOCATION: (0)...(0)
OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-24
Query Match 90.9%; Score 179; DB 3; Length 44;
Best Local Similarity 100.0%; Pred. No. 2.1e-23;
Matches 44; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DB 1 ANSFLXXLRGSLXRCIXXICDPXAKXIFEDVDTLAFWSKH 44
1 ANSFLXXLRGSLXRCIXXICDPXAKXIFEDVDTLAFWSKH 44
RESULT 2
US-10-182-263-6
Sequence 6, Application US/10182263
Patent No. 6530138
GENERAL INFORMATION:
APPLICANT: Gerlitz, Bruce E
APPLICANT: Jones, Bryan B
APPLICANT: Grinnell, Brian W
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13611
CURRENT APPLICATION NUMBER: US/10/182,263
CURRENT FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2002-02-11
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
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SEQ ID NO 6
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-182-263-6

Query Match 90.3%; Score 179; DB 4; Length 419;
Best Local Similarity 79.5%; Pred. No. 2; 8e-22;
Matches 35; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

Oy 1 ANSFLXXLRGSLRXCIXXICDPXXAKXIFEDVDDTLAFMSKH 44
Db 1 ANSFLXELRGSLERECIEICDFEAKXIFEDVDDTLAFMSKH 44

RESULT 3
US-08-955-636-35
Sequence 35, Application US/08955636A
Patent No. 6017882
GENERAL INFORMATION:
APPLICANT: Nelsentuen, Gary
TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
FILE REFERENCE: 09531/002001
CURRENT FILING DATE: 1997-10-23
CURRENT APPLICATION NUMBER: US/08/955,636A
NUMBER OF SEQ ID NOS: 35
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 35
LENGTH: 44
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: MOD_RES
LOCATION: (0)...(0)
OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-35

Query Match 89.3%; Score 176; DB 3; Length 44;
Best Local Similarity 97.7%; Pred. No. 6; 9e-23;
Matches 43; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Oy 1 ANSFLXXLRGSLRXCIXXICDPXXAKXIFEDVDDTLAFMSKH 44
Db 1 ANSFLXXLRGSLRXCIXXICDPXXAKXIFEDVDDTLAFMSKH 44

RESULT 4
US-10-182-263-3
Sequence 3, Application US/10182263
Patent No. 6630138
GENERAL INFORMATION:
APPLICANT: Gerlitz, Bruce E
APPLICANT: Jones, Bryan E
APPLICANT: Grinnell, Brian W
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13611
CURRENT APPLICATION NUMBER: US/10/182,263
CURRENT FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2002-02-11
PRIOR APPLICATION NUMBER: 60/189199
PRIOR FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 3
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-182-263-3

Query Match 88.3%; Score 174; DB 4; Length 419;
Best Local Similarity 77.3%; Pred. No. 2e-21;

Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
Oy 1 ANSFLXXLRGSLRXCIXXICDPXXAKXIFEDVDDTLAFMSKH 44
Db 1 ANSFLXELRGSLERECIEICDFEAKXIFEDVDDTLAFMSKH 44

RESULT 5
US-10-182-263-4
Sequence 4, Application US/10182263
Patent No. 6630138
GENERAL INFORMATION:
APPLICANT: Gerlitz, Bruce E
APPLICANT: Jones, Bryan E
APPLICANT: Grinnell, Brian W
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13611
CURRENT APPLICATION NUMBER: US/10/182,263
CURRENT FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2002-02-11
PRIOR APPLICATION NUMBER: 60/189199
PRIOR FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 4
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-182-263-4

Query Match 88.3%; Score 174; DB 4; Length 419;
Best Local Similarity 77.3%; Pred. No. 2e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Oy 1 ANSFLXXLRGSLRXCIXXICDPXXAKXIFEDVDDTLAFMSKH 44
Db 1 ANSFLXELRGSLERECIEICDFEAKXIFEDVDDTLAFMSKH 44

RESULT 6
US-10-182-263-5
Sequence 5, Application US/10182263
Patent No. 6630138
GENERAL INFORMATION:
APPLICANT: Gerlitz, Bruce E
APPLICANT: Jones, Bryan E
APPLICANT: Grinnell, Brian W
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13611
CURRENT APPLICATION NUMBER: US/10/182,263
CURRENT FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2002-02-11
PRIOR APPLICATION NUMBER: 60/189199
PRIOR FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 5
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-182-263-5

Query Match 88.3%; Score 174; DB 4; Length 419;
Best Local Similarity 77.3%; Pred. No. 2e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Oy 1 ANSFLXXLRGSLRXCIXXICDPXXAKXIFEDVDDTLAFMSKH 44
Db 1 ANSFLXELRGSLERECIEICDFEAKXIFEDVDDTLAFMSKH 44

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RESULT 7
US-08-955-636-20
; Sequence 20, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelstuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; TITLE OF INVENTION: POLYPEPTIDES
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 20
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (0)...(0)
; OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-20
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Query Match      87.8%; Score 173; DB 3; Length 44;
Best Local Similarity 97.7%; Pred. No. 2,2e-22;
Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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OY 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
Db 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
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RESULT 8
US-08-955-636-21
; Sequence 21, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelstuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; TITLE OF INVENTION: POLYPEPTIDES
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 21
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (0)...(0)
; OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-21
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Query Match      86.3%; Score 170; DB 3; Length 44;
Best Local Similarity 95.5%; Pred. No. 7,2e-22;
Matches 42; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
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```
OY 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
Db 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
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RESULT 9
US-08-955-636-19
; Sequence 19, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelstuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; TITLE OF INVENTION: POLYPEPTIDES
; FILE REFERENCE: 09531/002001
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; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 19
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (0)...(0)
; OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-19
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Query Match      85.3%; Score 168; DB 3; Length 44;
Best Local Similarity 95.5%; Pred. No. 1,6e-21;
Matches 42; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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OY 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
Db 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
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RESULT 10
US-08-955-636-22
; Sequence 22, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelstuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; TITLE OF INVENTION: POLYPEPTIDES
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 22
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (0)...(0)
; OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-22
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Query Match      85.3%; Score 168; DB 3; Length 44;
Best Local Similarity 95.5%; Pred. No. 1,6e-21;
Matches 42; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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OY 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
Db 1 ANSFLXLRQGSILRXKXICIXICDPXXAKXIFEDVDVDTLAFWSKH 44
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RESULT 11
US-08-955-636-1
; Sequence 1, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelstuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; TITLE OF INVENTION: POLYPEPTIDES
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
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NAME/KEY: MOD_RES
LOCATION: (0)...(0)
OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-1

Query Match 81.2%; Score 160; DB 3; Length 44;
Best Local Similarity 90.9%; Pred. No. 3.6e-20;
Matches 40; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 ANSFLXXLRSGSLXRXCIXXICDPXXAKXIFEDVDPTLAFWSKH 44
DB 1 ANSFLXXLRHSSSLXRXCIXXICDPXXAKXIFQVVDPTLAFWSKH 44

RESULT 12
US-08-955-636-25
Sequence 25, Application US/08955636A
Patent No. 6017882
GENERAL INFORMATION:
APPLICANT: Neisester, Gary
TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
FILE REFERENCE: 09531/002001
CURRENT APPLICATION NUMBER: US/08/955,636A
CURRENT FILING DATE: 1997-10-23
NUMBER OF SEQ ID NOS: 35
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 25
LENGTH: 44
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: MOD RES
LOCATION: (0)...(0)
OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid
US-08-955-636-25

Query Match 81.2%; Score 160; DB 3; Length 44;
Best Local Similarity 93.2%; Pred. No. 3.6e-20;
Matches 41; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 ANSFLXXLRSGSLXRXCIXXICDPXXAKXIFEDVDPTLAFWSKH 44
DB 1 ANSFLXXLRHSSSLXRXCIXXICDPXXAKXIFEDVDPTLAFWSKH 44

RESULT 13
US-08-965-832-2
Sequence 2, Application US/08965832
Patent No. 5847085
GENERAL INFORMATION:
APPLICANT: CHARLES T. ESMON AND MIKHAIL D. SMIRNOV
TITLE OF INVENTION: Modified Protein C
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Patrea L. Pabst
STREET: 2800 One Atlantic Center, 1201 West
STREET: Peachtree Street
CITY: Atlanta
STATE: GA
COUNTRY: USA
ZIP: 30309-3450
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/965,832
FILING DATE: 7-NOV-1997
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/745,254

FILING DATE: 8-NOV-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/053,768
FILING DATE: 25-JUL-1997
ATTORNEY/AGENT INFORMATION:
NAME: Pabst, Patrea L.
REGISTRATION NUMBER: 31,284
REFERENCE/DOCKET NUMBER: OMRP 165/167
TELECOMMUNICATION INFORMATION:
TELEPHONE: (404)-873-8794
TELEFAX: (404)-873-8795

INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 45 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear

MOLECULE TYPE: peptide
FEATURE:
NAME/KEY:
LOCATION: 6, 7, 14, 16, 19, 20, 25, 26, 29
OTHER INFORMATION: /note= "where Xaa means gamma
OTHER INFORMATION: carboxyglutamic acid"
FEATURE:
NAME/KEY:
LOCATION:
OTHER INFORMATION: /note= "partial sequence of human protein C"

US-08-965-832-2
Query Match 81.2%; Score 160; DB 2; Length 45;
Best Local Similarity 90.9%; Pred. No. 3.7e-20;
Matches 40; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 ANSFLXXLRSGSLXRXCIXXICDPXXAKXIFEDVDPTLAFWSKH 44
DB 1 ANSFLXXLRHSSSLXRXCIXXICDPXXAKXIFQVVDPTLAFWSKH 44

RESULT 14
US-08-295-411-1
Sequence 1, Application US/08295411
Patent No. 5679639
GENERAL INFORMATION:
APPLICANT: Griffin, John H.
APPLICANT: Meisters, Rolf M.
TITLE OF INVENTION: Serine protease-Derived Polypeptides and
TITLE OF INVENTION: Anti-Peptide Antibodies Systems and Therapeutic Methods
TITLE OF INVENTION: For Inhibiting Coagulation
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Office of Patent Counsel, The Scripps
ADDRESSEE: Research Institute
STREET: 10666 No. 5679639th Torrey Pines Road, TPC 8
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/295,411
FILING DATE: 22-AUG-1994
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/793,989
FILING DATE: 18-NOV-1991
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: Filling, Thomas
REGISTRATION NUMBER: 34,163

REFERENCE/DOCKET NUMBER: TSRI263.0C1
TELEPHONE: 619-554-2937
TELEFAX: 619-554-2937
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 419 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: Region
LOCATION: 1..157
OTHER INFORMATION: /note= "Protein C Light Chain"
FEATURE:
NAME/KEY: Region
LOCATION: 158..169
OTHER INFORMATION: /note= "Protein C Activation"
OTHER INFORMATION: Peptide
FEATURE:
NAME/KEY: Region
LOCATION: 170..419
OTHER INFORMATION: /note= "Protein C Heavy Chain"
US-08-295-411-1
Query Match
Best Local Similarity 81.2%; Score 160; DB 1; Length 419;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;
Qy 1 ANSFLXLRQSLKXKICIXICDPFXAKXIFEDVDTLAFWSKH 44
Db 1 ANSFLBELRHSLERECIEICDFEAKXIFQNVDDTLAFWSKH 44
RESULT 15
US-08-955-471-1
Sequence 1, Application US/08955471
Patent No. 5968751
GENERAL INFORMATION:
APPLICANT: Griffin, John H.
APPLICANT: Mesters, Rolf M.
TITLE OF INVENTION: Serine Protease-Derived Polypeptides and
Anti-Peptide Antibodies, Systems and Therapeutic Methods
TITLE OF INVENTION: for Inhibiting Coagulation
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Office of Patent Counsel, The Scripps
Research Institute
STREET: 10666 No. 5968751th Torrey Pines Road, TPC 8
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/955,471
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/295,411
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Fitting, Thomas
REGISTRATION NUMBER: 34,163
REFERENCE/DOCKET NUMBER: TSRI263.0C1
TELECOMMUNICATION INFORMATION:

TELEPHONE: 619-554-2937
TELEFAX: 619-554-2937
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 419 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: Region
LOCATION: 1..157
OTHER INFORMATION: /note= "Protein C Light Chain"
FEATURE:
NAME/KEY: Region
LOCATION: 158..169
OTHER INFORMATION: /note= "Protein C Activation"
OTHER INFORMATION: Peptide
FEATURE:
NAME/KEY: Region
LOCATION: 170..419
OTHER INFORMATION: /note= "Protein C Heavy Chain"
US-08-955-471-1
Query Match
Best Local Similarity 81.2%; Score 160; DB 2; Length 419;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;
Qy 1 ANSFLXLRQSLKXKICIXICDPFXAKXIFEDVDTLAFWSKH 44
Db 1 ANSFLBELRHSLERECIEICDFEAKXIFQNVDDTLAFWSKH 44
Search completed: March 1, 2004, 10:12:21
Job time : 17.5 secs

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OM protein - protein search, using sw model

Run on: March 1, 2004, 09:55:12 ; Search time 37.5 Seconds
(without alignments)
370.208 Million cell updates/sec

Title: SEQ1-4SUB5
Perfect score: 197
Sequence: 1 ANSFLXXLRGSLXKXIX.....XXAKXIFEDVDLAFWSKH 44

Scoring table: ELOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

1: SP_ARCHAEA:*
2: SP_BACTERIA:*
3: SP_FUNGI:*
4: SP_HUMAN:*
5: SP_INVERTEBRATE:*
6: SP_MAMMAL:*
7: SP_MNC:*
8: SP_ORNITHINE:*
9: SP_PHAGE:*
10: SP_PLANT:*
11: SP RODENT:*
12: SP_VIRUS:*
13: SP_VERTEBRATE:*
14: SP_UNCLASSIFIED:*
15: SP_VIRUS:*
16: SP_BACTERIAP:*
17: SP_ARCHAEP:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	151	76.6	456	6	Q9TRR0
2	140	71.1	460	11	Q91WN8
3	134	68.0	460	11	Q99PC6
4	118	59.9	55	4	Q8J0C2
5	118	59.9	55	4	Q8J0C5
6	115	58.4	482	11	Q63207
7	109	55.3	455	13	Q7SY86
8	107	54.3	433	13	Q8N2N6
9	107	54.3	433	13	Q8O4X5
10	105	52.5	524	13	Q7EXH8
11	103.5	51.3	443	13	Q8JHC9
12	101	51.3	340	11	Q80Y26
13	101	51.3	481	11	Q54740
14	101	51.3	481	11	Q98132
15	101	51.3	481	11	Q88947
16	99.5	50.5	442	13	Q804X1

17	99	50.3	679	4	Q96PQ8	Q96PQ8 homo sapien
18	97	49.2	474	13	Q8JHC8	Q8JHC8 brachydanio
19	95	48.2	469	6	Q9GMD9	Q9GMD9 onitrotrypn
20	93	47.2	229	13	Q8J040	Q8J040 xenopus lae
21	93	47.2	434	13	Q713B6	Q713B6 brachydanio
22	92	46.7	268	4	Q8NEK6	Q8NEK6 homo sapien
23	87	44.2	425	13	Q804X7	Q804X7 gallus gall
24	87	44.2	612	13	Q804W7	Q804W7 fugu rubrip
25	85	43.1	475	13	Q804W9	Q804W9 fugu rubrip
26	85	43.1	497	4	Q72715	Q72715 homo sapien
27	85	43.1	650	4	Q16519	Q16519 homo sapien
28	85	43.1	650	4	Q9NSD0	Q9NSD0 homo sapien
29	84	42.6	100	4	Q15253	Q15253 homo sapien
30	84	42.6	622	4	Q727P3	Q727P3 homo sapien
31	82.5	41.9	542	5	Q8T613	Q8T613 halocynthia
32	81	41.1	471	13	Q804X6	Q804X6 gallus gall
33	80	40.6	446	11	Q61109	Q61109 mus musculu
34	80	40.6	461	6	Q95ND7	Q95ND7 pan troglod
35	80	40.6	461	6	Q95ND6	Q95ND6 pan troglod
36	79	40.1	441	13	Q804X2	Q804X2 fugu rubrip
37	78	39.6	138	6	Q28994	Q28994 sus scrofa
38	78	39.6	607	13	Q91001	Q91001 gallus gall
39	78	39.6	648	6	Q29094	Q29094 sus scrofa
40	77	39.1	52	4	Q81XD5	Q81XD5 homo sapien
41	76	38.6	446	11	Q8K3U6	Q8K3U6 rattus norv
42	74	37.6	52	4	Q81XC5	Q81XC5 homo sapien
43	73.5	37.3	433	13	Q90YK1	Q90YK1 brachydanio
44	73.5	37.3	433	13	Q90YK1	Q90YK1 brachydanio
45	73	37.1	49	6	Q95ME8	Q95ME8 bos taurus

ALIGNMENTS

RESULT 1

Q9TRR0 PRELIMINARY; PRT; 456 AA.
AC Q9TRR0;
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Protein C precursor.
GN PROC.
OS Canis familiaris (Dog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Euteria; Carnivora; Fissipedia; Canidae; Canis.
OX NCBI_TaxID=9615;
RN [1]
RP SEQUENCE FROM N.A.
RA Leeb T., Kopp T., Deppe A., Breen M., Natis U., Brunberg L.,
RT "Molecular characterization and chromosomal assignment of the canine
RT protein C gene";
RL Mamm. Genome 10:135-139(1999).
RU [2]
RN SEQUENCE FROM N.A.
RX MEDLINE=93371952; PubMed=10443005;
RA Leeb T., Pfeiffer I., Kopp T., Deppe A., Breen B.,
RT "Analysis of canine protein C gene polymorphisms";
RL Anim. Genet. 30:237-238(1999).
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL: AF001979; CAA05126.1;
DR HSPF: P04070; IANT.
DR GO: GO:0005576; Cytosolic; IEA.
DR GO: GO:0005509; Fcalcium ion binding; IEA.
DR GO: GO:0004263; Fcatalytic activity; IEA.
DR GO: GO:0008233; Fpeptidase activity; IEA.
DR GO: GO:0004295; Ftrypsin activity; IEA.
DR GO: GO:0006508; Fproteolysis and peptidolysis; IEA.
DR InterPro: IPR000152; Asx_hydroxyl_S.
DR InterPro: IPR009003; Cys_ser_trypsin.
DR InterPro: IPR01881; EGF_Ca.
DR InterPro: IPR006209; EGF_like.

DR InterPro; IPR002383; GLA_blood.
 DR InterPro; IPR006210; IEGF.
 DR InterPro; IPR001254; Peptidase_S1.
 DR InterPro; IPR001314; Peptidase_S1A.
 DR InterPro; IPR000294; VitK_dep_GLA.
 DR Pfam; PFO00089; EGF_2.
 DR Pfam; PFO0594; GLA_1.
 DR PRINTS; PR00722; CHYMOTRYPSIN.
 DR PRINTS; PR00101; GLABLOOD.
 DR SMART; SM00181; EGF_2.
 DR SMART; SM00069; GLA_1.
 DR SMART; SM00020; TRYPSIN_DOM; 1.
 DR PROSITE; PS00010; ASX_HYDROXYL; 1.
 DR PROSITE; PS00022; EGF_1; 1.
 DR PROSITE; PS00186; EGF_2; 2.
 DR PROSITE; PS01187; EGF_CA; 1.
 DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
 DR PROSITE; PS00240; TRYPSIN_DOM; 1.
 DR PROSITE; PS00134; TRYPSIN_HIS; 1.
 DR PROSITE; PS00135; TRYPSIN_SER; 1.
 DR EGF-like domain; Hydrolase; Protease; Serine protease; Signal.
 FT SIGNAL 1 42
 FT CHAIN 43 192 POTENTIAL.
 FT CHAIN 193 194 PROTEIN C LIGHT CHAIN.
 FT CHAIN 195 456 PROTEIN C HEAVY CHAIN.
 SQ SEQUENCE 456 AA; 50813 MW; 7AD3A8C1C34559FF CRC64;

Query Match 76.6%; Score 151; DB 6; Length 456;
 Best Local Similarity 63.6%; Pred. No. 3.8e-17;
 Matches 28; Conservative 6; Mismatches 10; Indels 0; Gaps 0;

QY 1 ANSFLXLRGSLRXCIXXICDPFXAKXIFEDVDPLAFMSKH 44
 DB 43 ANSFLERMRGSLRXCMEICDPFBAKQIFQNVDTLAFMIKY 86

RESULT 2
 ID Q91WN8 PRELIMINARY; PRT; 460 AA.
 AC Q91WN8;
 DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Similar to protein C.
 GN PROC.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_Taxid=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RA Strausberg R.;
 RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
 DR EMBL; AC013896; AAH1896.1; -
 DR HSSP; P04070; 1AUT.
 DR MGD; MGI:97771; Proc.
 DR GO; GO:0005576; C:extracellular; IEA.
 DR GO; GO:0005509; F:calcium ion binding; IEA.
 DR GO; GO:0004263; F:chymotrypsin activity; IEA.
 DR GO; GO:0008233; F:peptidase activity; IEA.
 DR GO; GO:0004295; F:trypsin activity; IEA.
 DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
 DR InterPro; IPR000152; Asx_hydroxyl_S.
 DR InterPro; IPR009003; Cys_Ser_trypsin.
 DR InterPro; IPR001881; EGF_CA.
 DR InterPro; IPR006209; EGF_like.
 DR InterPro; IPR002383; GLA_blood.
 DR InterPro; IPR001254; Peptidase_S1.
 DR InterPro; IPR001314; Peptidase_S1A.
 DR InterPro; IPR000294; VitK_dep_GLA.

DR Pfam; PFO00089; EGF_2.
 DR Pfam; PFO0594; GLA_1.
 DR PRINTS; PR00722; CHYMOTRYPSIN.
 DR PRINTS; PR00001; GLABLOOD.
 DR SMART; SM00179; EGF_CA; 1.
 DR SMART; SM00069; GLA_1.
 DR SMART; SM00020; TRYPSIN_DOM; 1.
 DR PROSITE; PS00010; ASX_HYDROXYL; 1.
 DR PROSITE; PS00022; EGF_1; 1.
 DR PROSITE; PS01186; EGF_2; 2.
 DR PROSITE; PS01187; EGF_CA; 1.
 DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
 DR PROSITE; PS00240; TRYPSIN_DOM; 1.
 DR PROSITE; PS00134; TRYPSIN_HIS; 1.
 DR PROSITE; PS00135; TRYPSIN_SER; 1.
 DR EGF-like domain; Hydrolase; Protease; Serine protease.
 SQ SEQUENCE 460 AA; 51818 MW; 011F26E6FCC274 CRC64;

Query Match 71.1%; Score 140; DB 11; Length 460;
 Best Local Similarity 59.1%; Pred. No. 2.9e-15;
 Matches 26; Conservative 7; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXLRGSLRXCIXXICDPFXAKXIFEDVDPLAFMSKH 44
 DB 42 ANSFLERMRGSLRXCMEICDPFBAKQIFQNVDTLAFMIKY 85

RESULT 3
 ID Q99PC6 PRELIMINARY; PRT; 460 AA.
 AC Q99PC6;
 DT 01-JUN-2001 (TrEMBLrel. 17, Created)
 DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
 DE Anticoagulant protein C.
 GN PROC.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_Taxid=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/
 RA Kott I.;
 RT "Complete sequence of UC72A01."
 RL Submitted (NOV-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
 DR EMBL; AF318182; AK07918.1; -
 DR HSSP; P04070; 1AUT.
 DR MGD; MGI:97771; Proc.
 DR GO; GO:0005576; C:extracellular; IEA.
 DR GO; GO:0005509; F:calcium ion binding; IEA.
 DR GO; GO:0004263; F:chymotrypsin activity; IEA.
 DR GO; GO:0008233; F:peptidase activity; IEA.
 DR GO; GO:0004295; F:trypsin activity; IEA.
 DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
 DR InterPro; IPR000152; Asx_hydroxyl_S.
 DR InterPro; IPR009003; Cys_Ser_trypsin.
 DR InterPro; IPR001881; EGF_CA.
 DR InterPro; IPR006209; EGF_like.
 DR InterPro; IPR002383; GLA_blood.
 DR InterPro; IPR001254; Peptidase_S1.
 DR InterPro; IPR001314; Peptidase_S1A.
 DR InterPro; IPR000294; VitK_dep_GLA.
 DR Pfam; PFO00089; EGF_2.
 DR Pfam; PFO0594; GLA_1.
 DR PRINTS; PR00722; CHYMOTRYPSIN.
 DR PRINTS; PR00001; GLABLOOD.
 DR SMART; SM00179; EGF_CA; 1.
 DR SMART; SM00069; GLA_1.
 DR SMART; SM00020; TRYPSIN_DOM; 1.

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DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLUT_CARBOXYLATION; 1.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease.
SQ SEQUENCE 460 AA; 51784 MW; 0293BC25E9D3ED16 CRC64;

Query Match
Best Local Similarity 58.0%; Score 134; DB 1; Length 460;
Matches 25; Conservative 7; Mismatches 12; Indels 0; Gaps 0;

Qy 1 ANSFLXXLRQGSIXKXCIXXICDPFXKXIFEDVDTLAFWSKH 44
Db 42 ANSFLERHPGSLERECIEICDLEFAQEIFQNVDTLAFWKY 85

RESULT 4
Q8J002 PRELIMINARY; PRT; 55 AA.
AC Q8J002;
DT 01-MAR-2003 (TRENBLrel. 23, Created)
DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DE Protein C (Fragment).
GN PROC.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kinoshita S., Iida H., Inoue S., Watanabe K., Kurizhara M., Wada Y.,
RA "One M., Dongchon K., Hamasaki N.;"
RA "Gene Analysis of Anticoagulation Factors in Japanese Thrombotic
RA Patients: Genetic Background of Thrombophilia in Japan.;"
RT Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB083700; BAC21172.1;
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR InterPro; IPR002383; GLA_blood.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00594; gla; 1.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00069; GLA; 1.
FT NON_TER 1
FT NON_TER 55
SQ SEQUENCE 55 AA; 6527 MW; 4F89496534A78836 CRC64;

Query Match
Best Local Similarity 59.9%; Score 118; DB 4; Length 55;
Matches 24; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

Qy 1 ANSFLXXLRQGSIXKXCIXXICDPFXKXIFEDVDT 37
Db 19 ANSFLERHPGSLERECIEICDLEFAQEIFQNVDT 55

RESULT 5
Q8IXB5 PRELIMINARY; PRT; 55 AA.
AC Q8IXB5;
DT 01-MAR-2003 (TRENBLrel. 23, Created)
DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DE Protein C (Fragment).
GN PROC1.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

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CX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Hamasaki S., Kang D., Kinoshita S., Iida K., Inoue S., Watanabe K.,
RA Kurizhara M., Wada Y., Ono M.;"
RA "Gene Analysis of Anticoagulation Factors in Japanese Thrombotic
RA Patients: Genetic Background of Thrombophilia in Japan.;"
RT Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB086851; BAC53631.1;
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR InterPro; IPR002383; GLA_blood.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00594; gla; 1.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00069; GLA; 1.
FT NON_TER 1
FT NON_TER 55
SQ SEQUENCE 55 AA; 6475 MW; 3803696534BC9289 CRC64;

Query Match
Best Local Similarity 59.9%; Score 118; DB 4; Length 55;
Matches 24; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

Qy 1 ANSFLXXLRQGSIXKXCIXXICDPFXKXIFEDVDT 37
Db 19 ANSFLERHPGSLERECIEICDLEFAQEIFQNVDT 55

RESULT 6
Q63207 PRELIMINARY; PRT; 482 AA.
AC Q63207;
DT 01-NOV-1996 (TRENBLrel. 01, Created)
DT 01-NOV-1996 (TRENBLrel. 01, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DE Factor X.
GN Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=Sprague-Dawley;
RA MEDLINE=86093366; Pubmed=8578339;
RA Stanton C., Ross R.P., Hutson S., Wallin R.;"
RT "Evidence for competition between vitamin K-dependent clotting factors
RT for intracellular processing by the vitamin K-dependent gamma-
RT carboxylase.;"
RL Thromb. Res. 80:63-73(1995).
CC -11 SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL; X79807; CAA56202.1;
DR PIR; S49075; EXRT.
DR HSP; P00742; IXKA.
DR MEROPS; S01.216;
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0008233; F:peptidase activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR000152; Asx_hydroxyl_S.
DR InterPro; IPR009003; Cys_ser_trypsin.
DR InterPro; IPR000742; EGF_2.
DR InterPro; IPR001881; EGF_CA.
DR InterPro; IPR001438; EGF_II.
DR InterPro; IPR006209; EGF_II.
DR InterPro; IPR002383; GLA_blood.
DR InterPro; IPR001254; Peptidase_S1.
DR InterPro; IPR001314; Peptidase_S1A.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00594; gla; 1.

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DR pfam: PF00089; trypsin; 1.
 DR PRINTS; PR00022; CHYMOTRYPSIN.
 DR PRINTS; PR00010; EGFLOD.
 DR PRINTS; PR00001; GLABLOD.
 DR SMART; SM00179; EGF_CA; 1.
 DR SMART; SM00069; GLA; 1.
 DR SMART; SM00020; tryp_Spc; 1.
 DR PROSITE; PS00010; ASX HYDROXYL; 1.
 DR PROSITE; PS00022; EGF_1; 1.
 DR PROSITE; PS01186; EGF_2; 2.
 DR PROSITE; PS01187; EGF_CA; 1.
 DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
 DR PROSITE; PS00240; TRYPSIN_DOM; 1.
 DR PROSITE; PS00134; TRYPSIN_HIS; 1.
 DR PROSITE; PS00135; TRYPSIN_SER; 1.
 DR EGF-like domain; Hydrolase; Protease; Serine protease.
 SQ SEQUENCE 482 AA; 54265 MW; 0284678E3954A698 CRC64;

Query Match 58.4%; Score 115; DB 11; Length 482;
 Best Local Similarity 43.2%; Pred. No. 6e-11;
 Matches 19; Conservative 10; Mismatches 15; Indels 0; Gaps 0;

QY 1 ANSFLXKLROGSLXRCXICXICDPFXKXIFEDVDITLAFMSKH 44
 DB 41 ANSFEEIKKGNLERECVEICSFEEAREVEFEDNEKTEFNKXY 84

RESULT 7

ID Q7SY86 PRELIMINARY; PRT; 455 AA.
 AC Q7SY86;
 DT 01-OCT-2003 (T-EMBLrel. 25, Created)
 DT 01-OCT-2003 (T-EMBLrel. 25, Last sequence update)
 DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
 DE Hypothetical protein.
 OS Xenopus laevis (African clawed frog).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipridae; Pipidae;
 OC Xenopodidae; Xenopus.
 NC NCB1_Taxid=8355;

RP SEQUENCE FROM N.A.
 RC TISSUE=Whole;
 RA MEDLINE=22341132; PubMed=12454917;
 RA Klein S.L., Strauberg R.L., Wagner L., Pontius J., Clifton S.W.,
 RA Richardson P.;
 RT "Genetic and genomic tools for Xenopus research: The NIH Xenopus
 RT initiative";
 RL Dev. Dyn. 225:384-391(2002).
 RN 121
 RP SEQUENCE FROM N.A.

RC TISSUE=Whole;
 RA MEDLINE=22388257; PubMed=12477932;
 RA Strauberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins P.S., Wagner L., Schenker C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buettow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Siegelman M., Soares M.B., Donald M.P., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Uediri T.B., Toshiyuki S., Carrinchi P., Prange C.J.,
 RA Rana S.S., Loggellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
 RA Bebak S.A., McGowan P.J., McKernan K.U., Malek J.A., Gannarathne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.W., Gay L.J., Huilyk S.W.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Farley J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakeley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez R.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
 RA Krzyzanski M.I., Skalka U., Smalls D.E., Scherch A., Schein J.E.,
 RA Jones S.J., Maira M.A.,
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).

RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Whole;
 RA Klein S., Strauberg R.;
 RL Submitted (JUL-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL; BC054968; AAH94968.1; --
 DR Hypothetical protein.
 SQ SEQUENCE 455 AA; 51811 MW; 07C027EDDB495330 CRC64;

Query Match 55.3%; Score 109; DB 13; Length 455;
 Best Local Similarity 50.0%; Pred. No. 6e-10;
 Matches 22; Conservative 6; Mismatches 16; Indels 0; Gaps 0;

QY 1 ANSFLXKLROGSLXRCXICXICDPFXKXIFEDVDITLAFMSKH 44
 DB 49 ANFMELKPKGSLERECIEKCDFEAREIFETKEDITLNFANKY 92

RESULT 8

ID Q8N2N6 PRELIMINARY; PRT; 231 AA.
 AC Q8N2N6;
 DT 01-OCT-2002 (T-EMBLrel. 22, Created)
 DT 01-OCT-2002 (T-EMBLrel. 22, Last sequence update)
 DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
 DE Hypothetical protein FLJ50093.
 OS Homo sapiens (human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 NC NCB1_Taxid=9606;

RP SEQUENCE FROM N.A.
 RC TISSUE=Embryo;
 RA Itoigai T., Oda T., Nishikawa T., Hayashi K., Otsuki T., Sugiyama T.,
 RA Suzuki Y., Nagai K., Sugano S., Ishii S., Kawai-Hiro Y., Saito K.,
 RA Yamamoto J., Wakamatsu A., Nakamura Y., Kojima S., Naganari K.,
 RA Maehiro Y., Ono T., Okano K., Yoshikawa Y., Aotaka S., Sasaki N.,
 RA Hattori A., Okumura K., Iwayanagi T., Minomiyu K.;
 RT "NEDD human cDNA sequencing project";
 RL Submitted (MAR-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AK074574; BAC1069.1; --
 DR GO; GO:0005576; C:extracellular; IEA.
 DR GO; GO:0005509; F:calcium ion binding; IEA.
 DR InterPro; IPR002383; GLA_blood.
 DR InterPro; IPR00294; VitK_dep_GLA.
 DR Pfam; PF00594; gla; 1.
 DR PRINTS; PR00001; GLABLOOD.
 DR SMART; SM00069; GLA; 1.
 DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
 DR Hypothetical protein.
 KW SEQUENCE 231 AA; 25844 MW; 8A373B0D5C1D0D81 CRC64;

Query Match 54.3%; Score 107; DB 4; Length 231;
 Best Local Similarity 43.9%; Pred. No. 6.3e-10;
 Matches 18; Conservative 8; Mismatches 15; Indels 0; Gaps 0;

QY 1 ANSFLXKLROGSLXRCXICXICDPFXKXIFEDVDITLAFW 41
 DB 20 ANFLELRLROGTLERECMEICSYERKVEFENKKTWFEW 60

RESULT 9

ID Q804X5 PRELIMINARY; PRT; 433 AA.
 AC Q804X5;
 DT 01-JUN-2003 (T-EMBLrel. 24, Created)
 DT 01-JUN-2003 (T-EMBLrel. 24, Last sequence update)
 DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
 DE Anticoagulant protein C precursor (EC 3.4.21.69).
 NC NCB1_Taxid=9606;

OS Gallus gallus (Chicken).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;

OC Gallus.
 RX NCBI_TaxID=9031;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Davidson C.J., Hirt R.P., Lal K., Snell P., Elgar G.,
 RT "Comparative sequence analysis and molecular evolution of blood
 coagulation genes from Gallus gallus and Fugu rubripes."
 RL Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF465270; A033365.1; -
 DR GO: GO:0005576; C:extracellular; IEA.
 DR GO: GO:0005509; F:calcium ion binding; IEA.
 DR GO: GO:0004263; F:chymotrypsin activity; IEA.
 DR GO: GO:0016787; F:hydrolyase activity; IEA.
 DR GO: GO:0003808; F:protein C (activated) activity; IEA.
 DR GO: GO:0004285; F:trypsin activity; IEA.
 DR GO: GO:0006508; P:proteolysis and peptidolysis; IEA.
 DR InterPro: IPR000152; Asx hydroxyl S.
 DR InterPro: IPR009003; Cys Ser trypsin.
 DR InterPro: IPR000742; EGF_2.
 DR InterPro: IPR001881; EGF_Ca.
 DR InterPro: IPR006209; EGF_Like.
 DR InterPro: IPR002383; GLA blood.
 DR InterPro: IPR006210; IEGF.
 DR InterPro: IPR001254; peptidase S1.
 DR InterPro: IPR001314; peptidase S1A.
 DR InterPro: IPR000294; Vitr_dep_GLA.
 DR Pfam: PF00008; EGF_1.
 DR Pfam: PF00594; GLA_1.
 DR Pfam: PF00089; trypsin_1.
 DR PRINTS: PR00722; CHYMOTRYPSIN.
 DR PRINTS: PR00001; GLABLOOD.
 DR SMART: SM00181; EGF_2.
 DR SMART: SM00179; EGF_Ca_1.
 DR SMART: SM00069; GLA_1.
 DR SMART: SM00020; TRYP_SPC_1.
 DR PROSITE: PS00010; ASX_HYDROXYL_1.
 DR PROSITE: PS00022; EGF_1.
 DR PROSITE: PS01186; EGF_2_2.
 DR PROSITE: PS01187; EGF_Ca_1.
 DR PROSITE: PS00011; GLU_CARBOXYLATION_1.
 DR PROSITE: PS02040; TRYPSIN_DOM_1.
 DR PROSITE: PS00134; TRYPSIN_HIS_1.
 DR PROSITE: PS00135; TRYPSIN_SER_1.
 KW Hydrolyase.
 SQ SEQUENCE 433 AA; 4668 MW; E09DDE56D7DA23 CRC64;
 Query Match 54.3%; Score 107; DB 13; Length 433;
 Best Local Similarity 50.0%; Pred. No. 1, 2e-09;
 Matches 22; Conservative 5; Mismatches 17; Indels 0; Gaps 0;

OC
 QY 1 ANSFLXLRGSLKRCIXXICDPFXAKXIFEDVDPTLAFMSKH 44
 DB 40 ANSFLXLRGSLKRCIXXICDPFXAKXIFEDVDPTLAFMSKH 83
 RESULT 10
 Q7SKH8 PRELIMINARY; PRT; 524 AA.
 AC Q7SKH8;
 DT 01-OCT-2003 (Tremblrel. 25, Created)
 DT 01-OCT-2003 (Tremblrel. 25, Last sequence update)
 DT 01-OCT-2003 (Tremblrel. 25, Last annotation update)
 DE Hypothetical protein.
 OS Brachydanio rerio (Zebrafish) (Danio rerio).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Danio.
 OX NCBI_TaxID=7955;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC Tissue=Body;
 RX MEDLINE=22386257; PubMed=12477932;

RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Kaiser R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.P., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Wax S.I., Wang J., Hsieh P.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.W., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Uebachs T.B., Toshimaru S., Carninci P., Prange C.,
 RA Raha S.S., Loggiano N.A., Peters G.J., Adamson R.D., Mullany S.J.,
 RA Bosak S.A., McSwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Rhee S.J., Morley K.C., Hale S., Garcia A.M., Gay L.J., Hultey S.W.,
 RA Villion D.K., Mizny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Gehlert J., Helton E., Kerteman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shchepetnikov Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman U.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.W., Butterfield Y.S.,
 RA Kravitski M.I., Skalska U., Smalins D.E., Scherch A., Schein J.E.,
 RA Jones S.J., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human
 and mouse cDNA sequences."
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Body;
 RA Stausberg R.;
 RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL: BC055596; AAH55596.1; -
 KW Hypothetical protein.
 SQ SEQUENCE 524 AA; 59560 MW; 1B4E08119080325 CRC64;
 Query Match 53.3%; Score 105; DB 13; Length 524;
 Best Local Similarity 43.2%; Pred. No. 3, 4e-09;
 Matches 19; Conservative 6; Mismatches 17; Indels 0; Gaps 0;

OC
 QY 1 ANSFLXLRGSLKRCIXXICDPFXAKXIFEDVDPTLAFMSKH 44
 DB 42 ANSFLXLRGSLKRCIXXICDPFXAKXIFEDVDPTLAFMSKH 85
 RESULT 11
 Q8H9C9 PRELIMINARY; PRT; 443 AA.
 AC Q8H9C9;
 DT 01-OCT-2002 (Tremblrel. 22, Created)
 DT 01-OCT-2002 (Tremblrel. 22, Last sequence update)
 DT 01-OCT-2003 (Tremblrel. 25, Last annotation update)
 DE Coagulation factor VIII.
 OS Brachydanio rerio (Zebrafish) (Danio rerio).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Danio.
 OX NCBI_TaxID=7955;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Hannun-Haiah R., Day K., Jagadeeswaran P.;
 RT "Comprehensive analysis of blood coagulation pathways in Zebrafish:
 RT Evolution of coagulation factor genes and identification of zebrafish
 RT factor VIII."
 RL Blood Cells Mol. Dis. 0:0-0(2002).
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
 DR EMBL: AF519546; AAM88342.1; -
 DR EMBL: AF519546; AAM71000.1; -
 DR GO: GO:0005576; C:extracellular; IEA.
 DR GO: GO:0005509; F:calcium ion binding; IEA.
 DR GO: GO:0004263; F:chymotrypsin activity; IEA.
 DR GO: GO:0006233; F:peptidase activity; IEA.
 DR GO: GO:0004295; F:trypsin activity; IEA.
 DR GO: GO:0006508; P:proteolysis and peptidolysis; IEA.
 DR InterPro: IPR009003; Cys Ser trypsin.
 DR InterPro: IPR000742; EGF_2.
 DR InterPro: IPR001881; EGF_Ca.
 DR InterPro: IPR006209; EGF_Like.
 DR InterPro: IPR002383; GLA blood.
 DR InterPro: IPR006210; IEGF.

DR InterPro: IPR001254; Peptidase_S1.
DR InterPro: IPR001314; Peptidase_S1A.
DR InterPro: IPR000294; Vitk_dep_GLA.
DR Pfam: PF00008; EGF_1.
DR Pfam: PF00594; Gla; 1.
DR Pfam: PF00089; trypsin_1.
DR PRINTS: PR00722; CHYMOTRYPSIN.
DR PRINTS: PR00001; GLABLOOD.
DR SMART: SM00181; EGF_2.
DR SMART: SM00179; EGF_CA; 1.
DR SMART: SM00069; GLA; 1.
DR SMART: SM00020; Tryp_Spec; 1.
DR PROSITE: PS00022; EGF_1; 1.
DR PROSITE: PS01186; EGF_2; 2.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease.
KW SEQUENCE 443 AA; 48823 MW; 2D2504718AE94FF4 CRC64;
SQ

Query Match 52.5%; Score 103.5; DB 13; Length 443;
Best Local Similarity 47.6%; Pred. No. 5.1e-09;
Matches 20; Conservative 7; Mismatches 14; Indels 1; Gaps 1;
DB 1 ANS-FLXXLRQGSLSKXCIXXICDPFXAXXIFEDVDTLAFW 41
38 ANSGFLBEMKAGNLERCEVERICDYEAREVFEEDDRTKQFW 79

RESULT 12
ID Q80Y26 PRELIMINARY; PRT; 340 AA.
AC Q80Y26;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE F10 protein.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Embryo;
RX MEDLINE=22386257; PubMed=12477932;
RA Strausberg R.J., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Datchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stadelson M., Soares M.B., Bonaldo M.F., Cassavant T.L., Scheetz T.E.,
RA Brownstein W.J., Ueda T.B., Toshiyuki S., Carninci P., Pangue C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Millar S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hultix S.W.,
RA Villalón D.K., Wuzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Rodriguez A.C., Grimwood J.W., Green E.D., Dickson M.C.,
RA Krzywinski M.J., Skalska U., Smalins D.E., Schnerch A., Schein J.E.,
RA Jones S.J., Marra M.A.,
RT "Generation and initial analysis of more than 15,000 full-length human
and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Embryo;
RA Strausberg R.J.,
RL Submitted (APR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL: BC050219; AAH50219.1;
DR GO: GO:0005575; Cytoplasmic; IEA.
DR GO: GO:0005509; F:calcium ion binding; IEA.

DR GO: GO:0004295; P:trypsin activity; IEA.
DR GO: GO:0005508; P:proteolysis and peptidolysis; IEA.
DR InterPro: IPR00152; Asx_hydroxyl_S.
DR InterPro: IPR009003; Cys_ser_trypsin.
DR InterPro: IPR00742; EGF_2.
DR InterPro: IPR00181; EGF_CA.
DR InterPro: IPR001438; EGF_II.
DR InterPro: IPR006209; EGF_like.
DR InterPro: IPR002383; GLA blood.
DR InterPro: IPR006210; IEGF.
DR InterPro: IPR001254; Peptidase_S1.
DR InterPro: IPR000294; Vitk_dep_GLA.
DR Pfam: PF00008; EGF_1.
DR Pfam: PF00594; Gla; 1.
DR PRINTS: PR00010; EGFBLLOOD.
DR PRINTS: PR00001; GLABLOOD.
DR SMART: SM00181; EGF_2.
DR SMART: SM00179; EGF_CA; 1.
DR SMART: SM00069; GLA; 1.
DR PROSITE: PS00010; ASX_HYDROXYL; 1.
DR PROSITE: PS00022; EGF_1; 1.
DR PROSITE: PS01186; EGF_2; 2.
DR PROSITE: PS01187; EGF_CA; 1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
SQ SEQUENCE 340 AA; 38359 MW; EE252D6157720811 CRC64;
Query Match 51.3%; Score 101; DB 11; Length 340;
Best Local Similarity 38.6%; Pred. No. 1e-08;
Matches 17; Conservative 10; Mismatches 17; Indels 0; Gaps 0;

DB 1 ANS-FLXXLRQGSLSKXCIXXICDPFXAXXIFEDVDTLAFW 44
53 ANSGFLBEMKAGNLERCEVERICDYEAREVFEEDDRTKQFW 96
RESULT 13
ID O54740 PRELIMINARY; PRT; 481 AA.
AC O54740;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Coagulation factor X precursor (EC 3.4.21.6).
GN F10 OR F10.
OS Mus musculus (Mouse).
OC Plasmid phagescript.
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RX MEDLINE=98454993; PubMed=9793672;
RA Heidemann H.R., Kornemann R.E.,
RT "Cloning and recombinant expression of mouse coagulation factor X."
RL Thromb. Res. 92:33-41(1998).
CC -1. SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL: AJ222677; CAA10933.1;
DR HSSP: P00742; 1XKA.
DR MEROPS: S01_216; -.
DR MCD: MCD:103107; F10.
DR GO: GO:0005575; Cytoplasmic; IEA.
DR GO: GO:0046821; C:extrachromosomal DNA; IEA.
DR GO: GO:0003804; F:bioid coagulation factor X activity; IEA.
DR GO: GO:0005509; F:calcium ion binding; IEA.
DR GO: GO:0004263; F:chymotrypsin activity; IEA.
DR GO: GO:0008233; F:peptidase activity; IEA.
DR GO: GO:0004295; P:trypsin activity; IEA.
DR GO: GO:0005508; P:proteolysis and peptidolysis; IEA.
DR InterPro: IPR000152; Asx_hydroxyl_S.
DR InterPro: IPR009003; Cys_ser_trypsin.

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DR InterPro: IPR000742; EGF_2.
DR InterPro: IPR001881; EGF_Ca.
DR InterPro: IPR001438; EGF_11.
DR InterPro: IPR006209; EGF_like.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Peptidase_S1.
DR InterPro: IPR001314; Peptidase_S1A.
DR InterPro: IPR000294; VtK_dep_GLA.
DR Pfam: PF000083; trypsin_1.
DR Pfam: PF00594; Gla_1.
DR Pfam: PF00089; trypsin_1.
DR PRINTS: PR000722; CHYMOTRYPSIN.
DR PRINTS: PR00010; EGFLOOD.
DR SMART: SM00179; EGF_CA_1.
DR SMART: SM00069; GLA_1.
DR SMART: SM00020; TRYP_SPE; 1.
DR PROSITE: PS00010; ASX_HYDROXYL; 1.
DR PROSITE: PS00022; EGF_1; 1.
DR PROSITE: PS01186; EGF_2; 2.
DR PROSITE: PS01187; EGF_CA_1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease; Signal;
KW plasmid.
KM
FT SIGNAL.
FT CHAIN 1 40 POTENTIAL.
SQ SEQUENCE 481 AA; 53986 MW; CF702DESEPD97AE CRC64;

Query Match 51.3%; Score 101; DB 11; Length 481;
Best Local Similarity 38.6%; Pred. No. 1.5e-08;
Matches 17; Conservative 10; Mismatches 17; Indels 0; Gaps 0;

CY 1 ANSFLXLRQGSIXKXICDPFXKXKXIFEDVDTLAFWSKH 44
DB 41 ANSFPEFKKGNLEKCEMERICSYEVRNRIFFDEKTKRWTKY 84

RESULT 14
ID Q99L32 PRELIMINARY; PRT; 481 AA.
AC Q99L32;
DT 01-JUN-2001 (Tremblrel. 17, Created)
DT 01-JUN-2001 (Tremblrel. 17, Last sequence update)
DT 01-OCT-2003 (Tremblrel. 25, Last annotation update)
DE Coagulation factor X.
GN F10.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_Taxid=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Strausberg R.;
RT Submitted (Feb-2001) to the EMBL/GenBank/DBJ databases.
CC 1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL: BC003877; AA03877.1; -.
DR HSSP: P00742; 1XKA.
DR MEROPS: S01.216; -.
DR MGD; MGI:103107; F10.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0008233; F:peptidase activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro: IPR000152; Asx_hydroxyl_S
DR InterPro: IPR008003; Cys_Ser_trypsin.
DR InterPro: IPR000742; EGF_2.
DR InterPro: IPR001881; EGF_Ca.
DR InterPro: IPR001438; EGF_11.
DR InterPro: IPR006209; EGF_like.
DR InterPro: IPR001254; Peptidase_S1.
DR InterPro: IPR001314; Peptidase_S1A.
DR InterPro: IPR000294; VtK_dep_GLA.
DR Pfam: PF000083; trypsin_1.
DR Pfam: PF00594; Gla_1.
DR Pfam: PF00089; trypsin_1.
DR PRINTS: PR000722; CHYMOTRYPSIN.
DR PRINTS: PR00010; EGFLOOD.
DR SMART: SM00179; EGF_CA_1.
DR SMART: SM00069; GLA_1.
DR SMART: SM00020; TRYP_SPE; 1.
DR PROSITE: PS00010; ASX_HYDROXYL; 1.
DR PROSITE: PS00022; EGF_1; 1.
DR PROSITE: PS01186; EGF_2; 2.
DR PROSITE: PS01187; EGF_CA_1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease.
SQ SEQUENCE 481 AA; 54004 MW; BD88E96C8A0B7E7F CRC64;
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DR InterPro: IPR006209; EGF_like.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Peptidase_S1.
DR InterPro: IPR001314; Peptidase_S1A.
DR InterPro: IPR000294; VtK_dep_GLA.
DR Pfam: PF000083; trypsin_1.
DR Pfam: PF00594; Gla_1.
DR Pfam: PF00089; trypsin_1.
DR PRINTS: PR000722; CHYMOTRYPSIN.
DR PRINTS: PR00010; EGFLOOD.
DR SMART: SM00179; EGF_CA_1.
DR SMART: SM00069; GLA_1.
DR SMART: SM00020; TRYP_SPE; 1.
DR PROSITE: PS00010; ASX_HYDROXYL; 1.
DR PROSITE: PS00022; EGF_1; 1.
DR PROSITE: PS01186; EGF_2; 2.
DR PROSITE: PS01187; EGF_CA_1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease.
SQ SEQUENCE 481 AA; 54004 MW; BD88E96C8A0B7E7F CRC64;

Query Match 51.3%; Score 101; DB 11; Length 481;
Best Local Similarity 38.6%; Pred. No. 1.5e-08;
Matches 17; Conservative 10; Mismatches 17; Indels 0; Gaps 0;

CY 1 ANSFLXLRQGSIXKXICDPFXKXKXIFEDVDTLAFWSKH 44
DB 41 ANSFPEFKKGNLEKCEMERICSYEVRNRIFFDEKTKRWTKY 84

RESULT 15
ID O88947 PRELIMINARY; PRT; 481 AA.
AC O88947;
DT 01-NOV-1998 (Tremblrel. 08, Created)
DT 01-NOV-1998 (Tremblrel. 08, Last sequence update)
DT 01-OCT-2003 (Tremblrel. 25, Last annotation update)
DE Coagulation factor X precursor.
GN F10.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_Taxid=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN-C57BL/6 J. CBA. TISSUE: Liver;
RX MEDLINE: 96347933; PubMed: 9687791;
RA Liang Z., Cooper A., Deford M.E., Carmeliet P., Collen D.,
RA Castellino F.J., Rosen E.D.;
RT "Cloning and characterization of a cDNA encoding murine coagulation
RT factor X.";
RL Thromb. Haemost. 80:87-91(1998).
RN [2]
RP SEQUENCE FROM N.A.
RA Cooper A., Liang Z., Castellino F.J., Rosen E.D.;
RT "Cloning and characterization of the murine Factor X Gene.";
RL Thromb. Haemost. 010-012000;
CC 1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL: AF087644; AAC6345.1; -.
DR EMBL: AF211347; AAF2980.1; -.
DR HSSP: P00742; 1XKA.
DR MEROPS: S01.216; -.
DR MGD; MGI:103107; F10.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
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DR GO:0006508; F:proteolysis and peptidolysis; IEA.
DR InterPro; IPR000152; Asx_hydroxyl_S.
DR InterPro; IPR009003; Cys_Ser_trypsin.
DR InterPro; IPR000742; EGF_2.
DR InterPro; IPR001881; EGF_Ca.
DR InterPro; IPR001438; EGF_II.
DR InterPro; IPR006209; EGF_like.
DR InterPro; IPR002383; GLA_blood.
DR InterPro; IPR001254; Peptidase_S1.
DR InterPro; IPR001314; Peptidase_S1A.
DR InterPro; IPR00294; Vltk_dep_GLA.
DR Pfam; PF000594; Gla; 2.
DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00010; EGFBLOOD.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00178; EGF_CA; 1.
DR SMART; SM00069; GLA; 1.
DR SMART; SM00020; Tryp_Spc; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLUT_CARBOXYLATION; 1.
DR PROSITE; PS50240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease; Signal.
FT SIGNAL 1 40
FT CHAIN 1 481
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COAGULATION FACTOR X.
POTENTIAL.
Query Match 51.3%; Score 101; DB 11; Length 481;
Best Local Similarity 38.6%; Pred. NO. 1.5e-08;
Matches 17; Conservative 10; Mismatches 17; Indels 0; Gaps 0;
Qy 1 ANSPFLXLRQGSIXRXCIXICDFFXAXKIFEDVDITLAFWSKH 44
Db 41 ANSPFEPFKGNLRECEMEICSYEVVEIFEDDEKTKWTKY 84

Search completed: March 1, 2004, 10:02:52
Job time : 38.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 1, 2004, 10:01:28 ; Search time 28 Seconds
(without alignments)
331.812 Million cell updates/sec

Title: SEQ1-4SUBS
Perfect score: 197
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Gapop 10.0 , Gapext 0.5

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Minimum DB seq length: 0
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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	179	90.9	419	14	US-10-182-263-6
2	179	90.9	419	15	US-10-168-407-5
3	179	90.9	419	15	US-10-168-407-6
4	174	88.3	419	14	US-10-182-263-3
5	174	88.3	419	14	US-10-182-263-4
6	174	88.3	419	14	US-10-182-263-5
7	174	88.3	419	15	US-10-168-407-3
8	174	88.3	419	15	US-10-168-407-4
9	160	81.2	419	14	US-10-298-330-1
10	160	81.2	419	10	US-09-978-917A-4
11	160	81.2	419	14	US-10-182-263-1
12	160	81.2	419	15	US-10-168-407-1
13	160	81.2	461	10	US-08-978-917A-2
14	160	81.2	461	14	US-10-182-263-2
15	160	81.2	461	15	US-10-168-407-2

16	116	58.9	44	14	US-10-298-330-2	Sequence 2, Appl
17	110	55.8	139	15	US-10-360-101-232	Sequence 23, App
18	110	55.8	488	14	US-10-348-504-44	Sequence 44, Appl
19	110	55.8	488	14	US-10-407-123-27	Sequence 27, Appl
20	106	53.8	44	14	US-10-298-330-18	Sequence 18, Appl
21	99	50.3	42	16	US-10-038-854-97	Sequence 97, Appl
22	99	50.3	406	15	US-09-782-587B-3	Sequence 3, Appl
23	99	50.3	406	15	US-10-383-898-1	Sequence 1, Appl
24	99	50.3	466	14	US-10-017-122-2	Sequence 14, Appl
25	99	50.3	466	15	US-10-375-741-14	Sequence 14, Appl
26	96	48.7	44	14	US-10-298-330-3	Sequence 3, Appl
27	96	48.7	406	10	US-09-782-587B-1	Sequence 1, Appl
28	96	48.7	406	14	US-10-109-498-1	Sequence 1, Appl
29	96	48.7	406	14	US-10-255-032-1	Sequence 1, Appl
30	96	48.7	406	14	US-10-281-727-1	Sequence 1, Appl
31	96	48.7	406	15	US-10-386-898-7	Sequence 7, Appl
32	95	48.2	405	15	US-10-360-101-235	Sequence 225, Appl
33	91	46.2	40	14	US-10-298-330-23	Sequence 23, Appl
34	90	45.7	44	14	US-10-298-330-4	Sequence 23, Appl
35	89	45.2	40	14	US-10-298-330-25	Sequence 23, Appl
36	85	43.1	40	14	US-10-298-330-22	Sequence 22, Appl
37	84.5	42.9	96	10	US-09-759-130B-113	Sequence 313, App
38	84.5	42.9	96	14	US-10-189-123-43	Sequence 43, Appl
39	84.5	42.9	96	14	US-10-188-495-43	Sequence 43, Appl
40	84.5	42.9	209	10	US-09-759-130B-112	Sequence 312, App
41	84.5	42.9	209	14	US-10-189-123-42	Sequence 42, Appl
42	84.5	42.9	209	14	US-10-188-495-42	Sequence 42, Appl
43	84.5	42.9	226	10	US-09-759-130B-210	Sequence 310, App
44	84.5	42.9	226	14	US-10-189-123-40	Sequence 40, Appl
45	84.5	42.9	226	14	US-10-188-495-40	Sequence 40, Appl

ALIGNMENTS

RESULT 1
US-10-182-263-6
; Sequence 6, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian M
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT FILING DATE: 2002-07-22
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 6
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-6
Query Match 90.9%; Score 179; DB 14; Length 419;
Best Local Similarity 79.5%; Pred. No. 3.9e-22;
Matches 35; Conservative 0; Mismatches 9; Indels 0; Gaps 0;
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DB 1 ANSFLXLRGSLRXKXIXICPFXAKXIFEDVDTLAFWSKH 44
RESULT 2
US-10-168-407-5
; Sequence 5, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:

APPLICANT: Geriltz, Bruce E
APPLICANT: Jones, Bryan E
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13610
CURRENT APPLICATION NUMBER: US/10/168,407
CURRENT FILING DATE: 2002-11-04
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 5
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-168-407-5

Query Match 90.9%; Score 179; DB 15; Length 419;
Best Local Similarity 79.5%; Pred. No. 3.9e-22;
Matches 35; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXLRQGSIXKXCIXICDPFXKXIFEDVDITLAFWSKH 44
DB 1 ANSFLBELRQGSLERECEIEICDPFEAKXIFEDVDITLAFWSKH 44

RESULT 3
US-10-168-407-6
Sequence 6, Application US/10168407
Publication No. US20030207435A1
GENERAL INFORMATION:
APPLICANT: Geriltz, Bruce E
APPLICANT: Jones, Bryan E
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13610
CURRENT APPLICATION NUMBER: US/10/168,407
CURRENT FILING DATE: 2002-11-04
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 6
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-168-407-6

Query Match 90.9%; Score 179; DB 15; Length 419;
Best Local Similarity 79.5%; Pred. No. 3.9e-22;
Matches 35; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

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DB 1 ANSFLBELRQGSLERECEIEICDPFEAKXIFEDVDITLAFWSKH 44

RESULT 4
US-10-182-263-3
Sequence 3, Application US/10182263
Publication No. US20030022354A1
GENERAL INFORMATION:
APPLICANT: Geriltz, Bruce E
APPLICANT: Jones, Bryan E
APPLICANT: Grinnell, Brian W
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13611
CURRENT APPLICATION NUMBER: US/10/182,263
CURRENT FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2002-02-11
PRIOR APPLICATION NUMBER: 60/189199
PRIOR FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 3
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens

US-10-182-263-3

Query Match 88.3%; Score 174; DB 14; Length 419;
Best Local Similarity 77.3%; Pred. No. 2.8e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 ANSFLXLRQGSIXKXCIXICDPFXKXIFEDVDITLAFWSKH 44
DB 1 ANSFLBELRQGSLERECEIEICDPFEAKXIFEDVDITLAFWSKH 44

RESULT 5
US-10-182-263-4
Sequence 4, Application US/10182263
Publication No. US20030022354A1
GENERAL INFORMATION:
APPLICANT: Geriltz, Bruce E
APPLICANT: Jones, Bryan E
APPLICANT: Grinnell, Brian W
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13611
CURRENT APPLICATION NUMBER: US/10/182,263
CURRENT FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2002-02-11
PRIOR APPLICATION NUMBER: 60/189199
PRIOR FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 4
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-182-263-4

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Best Local Similarity 77.3%; Pred. No. 2.8e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

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DB 1 ANSFLBELRQGSLERECEIEICDPFEAKXIFEDVDITLAFWSKH 44

RESULT 6
US-10-182-263-5
Sequence 5, Application US/10182263
Publication No. US20030022354A1
GENERAL INFORMATION:
APPLICANT: Geriltz, Bruce E
APPLICANT: Jones, Bryan E
APPLICANT: Grinnell, Brian W
TITLE OF INVENTION: PROTEIN C DERIVATIVES
FILE REFERENCE: X-13611
CURRENT APPLICATION NUMBER: US/10/182,263
CURRENT FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: 60/181948
PRIOR FILING DATE: 2002-02-11
PRIOR APPLICATION NUMBER: 60/189199
PRIOR FILING DATE: 2000-03-14
NUMBER OF SEQ ID NOS: 12
SOFTWARE: PatentIn version 3.1
SEQ ID NO 5
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-10-182-263-5

Query Match 88.3%; Score 174; DB 14; Length 419;
Best Local Similarity 77.3%; Pred. No. 2.8e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

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Db 1 ANSFLELHSGSLERECIEICDFEAKEIFEDVDITLAFWSKH 44
RESULT 7
US-10-168-407-3
; Sequence 3, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 3
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-3

Query Match 88.3%; Score 174; DB 15; Length 419;
Best Local Similarity 77.3%; Pred. No. 2, 8e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 ANSFLLXLRQGSILXRCIXXICDFXXAKXIFEDVDITLAFWSKH 44
Db 1 ANSFLELHSGSLERECIEICDFEAKEIFEDVDITLAFWSKH 44

RESULT 8
US-10-168-407-4
; Sequence 4, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-4

Query Match 88.3%; Score 174; DB 15; Length 419;
Best Local Similarity 77.3%; Pred. No. 2, 8e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

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Db 1 ANSFLELHSGSLERECIEICDFEAKEIFEDVDITLAFWSKH 44

RESULT 9
US-10-298-330-1
; Sequence 1, Application US/10298330
; Publication No. US20030100506A1
; GENERAL INFORMATION:
; APPLICANT: Nelsestuen, Gary L.
; TITLE OF INVENTION: Modified Vitamin K-Dependent
; TITLE OF INVENTION: Polypeptides
; FILE REFERENCE: 09531-12700
; CURRENT APPLICATION NUMBER: US/10/298,330
; CURRENT FILING DATE: 2002-11-18
; PRIOR APPLICATION NUMBER: 09/497,591
; PRIOR FILING DATE: 2000-02-03

; PRIOR APPLICATION NUMBER: 09/302,239
; PRIOR FILING DATE: 1998-04-29
; PRIOR APPLICATION NUMBER: 08/955,636
; PRIOR FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 27
; SOFTWARE: FastSeq for Windows Version 4.0
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; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: VARIANT
; LOCATION: 6, 7, 14, 16, 19, 20, 25, 26, 29
; OTHER INFORMATION: Xaa = gamma carboxylutamic or glutamic acid
US-10-298-330-1

Query Match 81.2%; Score 160; DB 14; Length 44;
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Matches 40; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

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Db 1 ANSFLLXLRHSSILXRCIXXICDFXXAKXIFQVDDITLAFWSKH 44

RESULT 10
US-09-978-917A-4
; Sequence 4, Application US/09978917A
; Publication No. US20030027299A1
; GENERAL INFORMATION:
; APPLICANT: Maxygen Aps; Maxygen Holdings
; TITLE OF INVENTION: Protein C or activated protein C-like molecules
; FILE REFERENCE: 0219us310 - protein C
; CURRENT APPLICATION NUMBER: US/09/978,917A
; CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-978-917A-4

Query Match 81.2%; Score 160; DB 10; Length 419;
Best Local Similarity 70.5%; Pred. No. 7e-19;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

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Db 1 ANSFLELHSGSLERECIEICDFEAKEIFQVDDITLAFWSKH 44

RESULT 11
US-10-182-263-1
; Sequence 1, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 419
; TYPE: PRT

ORGANISM: Homo sapiens
US-10-182-263-1

Query Match 81.2%; Score 160; DB 14; Length 419;
Best Local Similarity 70.5%; Pred. No. 7e-19;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

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DB 1 ANSFLBELRHSSLERECIEICDPEAKKIFQNVDDTLAFWSKH 44

RESULT 12
US-10-168-407-1
; Sequence 1, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-1

Query Match 81.2%; Score 160; DB 15; Length 419;
Best Local Similarity 70.5%; Pred. No. 7e-19;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

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DB 1 ANSFLBELRHSSLERECIEICDPEAKKIFQNVDDTLAFWSKH 44

RESULT 13
US-09-978-917A-2
; Sequence 2, Application US/09978917A
; Publication No. US2003027299A1
; GENERAL INFORMATION:
; APPLICANT: Maxigen Aps; Maxigen Holdings
; TITLE OF INVENTION: Protein C or activated protein C-like molecules
; FILE REFERENCE: 0219us310 - protein C
; CURRENT APPLICATION NUMBER: US/09/978,917A
; CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 461
; TYPE: PRT
; ORGANISM: Homo sapiens
; NAME/KEY: SIGNAL
; LOCATION: (1)...(42)
; FEATURE:
; NAME/KEY: CHAIN
; LOCATION: (43)...(461)
US-09-978-917A-2

Query Match 81.2%; Score 160; DB 10; Length 461;
Best Local Similarity 70.5%; Pred. No. 7.8e-19;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

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DB 43 ANSFLBELRHSSLERECIEICDPEAKKIFQNVDDTLAFWSKH 86

RESULT 14
US-10-182-263-2
; Sequence 2, Application US/10182263
; Publication No. US2003022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 461
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-2

Query Match 81.2%; Score 160; DB 14; Length 461;
Best Local Similarity 70.5%; Pred. No. 7.8e-19;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXXLRGSLRXKXICXIXXICDPFXAXKXIFEDVDVDTLAFWSKH 44
DB 43 ANSFLBELRHSSLERECIEICDPEAKKIFQNVDDTLAFWSKH 86

RESULT 15
US-10-168-407-2
; Sequence 2, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 461
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-2

Query Match 81.2%; Score 160; DB 15; Length 461;
Best Local Similarity 70.5%; Pred. No. 7.8e-19;
Matches 31; Conservative 2; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXXLRGSLRXKXICXIXXICDPFXAXKXIFEDVDVDTLAFWSKH 44
DB 43 ANSFLBELRHSSLERECIEICDPEAKKIFQNVDDTLAFWSKH 86

Search completed: March 1, 2004, 10:11:10
Job time : 28 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 1, 2004, 09:54:37 ; Search time 50 Seconds
(without alignments)
248.642 Million cell updates/sec

Title: SEQ1-32GLU-33ASP
Perfect score: 198
Sequence: 1 ANSFLXLRHSLXRCIX.....XXAKXIFEDVDTAFMSKH 44

Scoring table: BLOSUM62
Gapop 10.0, Gapept 0.5

Searched: 1586107 seqs, 282547505 residues
Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: A_Geneseq_29Jan04:*

- 1: geneseqp19808:*
- 2: geneseqp19908:*
- 3: geneseqp20008:*
- 4: geneseqp20018:*
- 5: geneseqp20028:*
- 6: geneseqp20038:*
- 7: geneseqp20048:*
- 8: geneseqp20058:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	180	90.9	44	2	AAV18297 Modified
2	176	88.9	419	4	AAH82676 Human pro
3	176	88.9	419	4	AAH82675 Human pro
4	176	88.9	419	4	AAH82678 Human pro
5	176	88.9	419	4	AAH82677 Human pro
6	176	88.9	419	4	AAH82679 Human pro
7	174	87.9	45	2	AAH82676 Human pro
8	172	86.9	44	2	AAH82675 Human pro
9	172	86.9	44	2	AAH82678 Human pro
10	172	86.9	44	2	AAH82677 Human pro
11	172	86.9	44	2	AAH82679 Human pro
12	172	86.9	44	2	AAH82676 Human pro
13	172	86.9	44	2	AAH82675 Human pro
14	172	86.9	44	2	AAH82678 Human pro
15	172	86.9	44	2	AAH82677 Human pro
16	172	86.9	44	2	AAH82679 Human pro
17	172	86.9	44	2	AAH82676 Human pro
18	172	86.9	44	2	AAH82675 Human pro
19	172	86.9	44	2	AAH82678 Human pro
20	172	86.9	44	2	AAH82677 Human pro
21	172	86.9	44	2	AAH82679 Human pro
22	172	86.9	44	2	AAH82676 Human pro
23	172	86.9	44	2	AAH82675 Human pro
24	172	86.9	44	2	AAH82678 Human pro
25	172	86.9	44	2	AAH82677 Human pro

26	172	86.9	419	5	AAU99005 Human pro
27	172	86.9	419	5	AAU99006 Human pro
28	172	86.9	419	5	AAU99008 Human pro
29	172	86.9	419	5	AAU99010 Human pro
30	172	86.9	419	5	AAU99012 Human pro
31	172	86.9	419	5	AAU99014 Human pro
32	172	86.9	419	5	AAU99016 Human pro
33	172	86.9	419	5	AAU99018 Human pro
34	172	86.9	419	5	AAU99020 Human pro
35	172	86.9	419	5	AAU99022 Human pro
36	172	86.9	419	5	AAU99024 Human pro
37	172	86.9	419	5	AAU99026 Human pro
38	172	86.9	419	5	AAU99028 Human pro
39	172	86.9	419	5	AAU99030 Human pro
40	172	86.9	419	5	AAU99032 Human pro
41	172	86.9	419	5	AAU99034 Human pro
42	172	86.9	419	5	AAU99036 Human pro
43	172	86.9	419	5	AAU99038 Human pro
44	172	86.9	419	5	AAU99040 Human pro
45	172	86.9	419	5	AAU99042 Human pro

ALIGNMENTS

RESULT 1

AAV18297 standard; peptide; 44 AA.

AAV18297;

17-AUG-1999 (first entry)

Modified GUA domain of vitamin K-dependent protein.

GUA domain; muten; vitamin K-dependent protein; clotting disorder;

therapy.

Homo sapiens.

Synthetic.

Key

Misc-difference 1..44

/note= "Xaa= gamma-carboxyglutamic acid, or glutamic acid"

W09920767-A1.

23-APR-1999.

20-OCT-1996; 98WO-US022152.

23-OCT-1997; 97US-00955636.

(MINT) UNIV MINNESOTA.

Nelsetuen GL;

WPI; 1999-288309/24.

Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid domain, useful for treating clotting disorders.

Claim 6, Page 78; 86PP; English.

This sequence represents a modified GUA (gamma-carboxyglutamic acid) domain. The invention relates to a vitamin K-dependent polypeptide

comprising a modified GUA domain containing an amino acid substitution

which enhances membrane binding of the modified polypeptide as compared

to the native polypeptide. The polypeptide is used to treat a clotting

disorder by decreasing or increasing clot formation. Modification of the

GUA domain results in a protein which has enhanced membrane binding

affinity as compared to the native protein

XX Sequence 44 AA;
 Query Match 90.9%; Score 180; DB 2; Length 44;
 Best Local Similarity 100.0%; Pred. No. 3,6e-22;
 Matches 44; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 ANSFLXLRHSSLRKXCIXICDPFXAKXIFEDVDPTLAFWSKH 44
 1 ANSFLXLRHSSLRKXCIXICDPFXAKXIFEDVDPTLAFWSKH 44

RESULT 2
 AAB82676
 ID AAB82676 standard; protein; 419 AA.
 AC AAB82676;
 XX
 DT 15-OCT-2001 (first entry)

Human protein C derivative (S11G/Q32E/N33D/L194S/T254S).
 Protein C; human; coronary syndrome; thrombosis; angina;
 myocardial infarction; vascular occlusive disorder; hypercoagulation;
 sepsis; protein C deficiency; occlusion; thromboembolism; stenosis;
 antibacterial; immunosuppressive; thrombolytic; cardiant; antianginal;
 anticoagulant; therapy; mutant; mutein.

OS Homo sapiens.
 OS Synthetic.
 FH Key
 FT Location/Qualifiers
 FT 1..45
 FT Domain
 FT /note= "Gla domain"
 FT 6
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 7
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 11
 FT Misc-difference
 FT /note= "Ser in wild-type protein"
 FT 14
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 16
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 19
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 20
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 25
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 26
 FT Modified-site
 FT /note= "gamma-carboxylated"
 FT 29
 FT Modified-site
 FT /note= "N-glycosylated"
 FT 32
 FT Misc-difference
 FT /note= "Gln in wild-type protein"
 FT 33
 FT Misc-difference
 FT /note= "Asn in wild-type protein"
 FT 50..69
 FT Disulfide-bond
 FT 58..64
 FT Disulfide-bond
 FT 80..89
 FT Disulfide-bond
 FT 98..109
 FT Disulfide-bond
 FT 120..133
 FT Disulfide-bond
 FT 141..277
 FT Disulfide-bond
 FT 156..157
 FT Cleavage-site
 FT /note= "cleavage makes a 2-chain inactive precursor (155-
 amino acid light chain attached via a disulfide bond to a
 262-amino acid heavy chain)"
 FT 158..168
 FT Peptide
 FT /note= "activation peptide; removal activates the 2-chain
 zymogen"
 FT 169..170
 FT Cleavage-site
 FT /note= "thrombin cleavage site"

FT Misc-difference 194
 FT /note= "Leu in wild-type protein"
 FT Disulfide-bond 196..212
 FT Modified-site 248
 FT /note= "N-glycosylated"
 FT Misc-difference 254
 FT /note= "Thr in wild-type protein"
 FT Modified-site 313
 FT /note= "N-glycosylated"
 FT Modified-site 329
 FT /note= "N-glycosylated"
 FT Disulfide-bond 332..345
 FT Disulfide-bond 356..384

WO200157193-A2.
 09-AUG-2001.
 19-JAN-2001; 2001WO-US000020.
 02-FEB-2000; 2000US-0179801P.
 14-MAR-2000; 2000US-0189197P.
 (ELIL) LILLY & CO ELI.
 Gerlitz BE; Jones BE;
 WPI; 2001-496919/54.
 N-PEDB; AAB82676.

Novel human protein C derivative for treating, e.g., myocardial
 infarction, unstable angina, sepsis, thrombotic disorders, acute arterial
 thrombotic occlusion, and thromboembolism.

Claim 4; Page 53-54; 63pp; English.

The present sequence is that of a claimed human protein C derivative in
 which Ser at position 11 of the mature wild-type protein C sequence (see
 AAB82673) is substituted with Gly, Gln at position 32 with Gln, Asn at
 position 33 with Asp, Leu at position 194 with Ser, and Thr at position
 254 with Ser. It is an example of protein C derivatives of the invention
 that have at least 2 amino acid substitutions, but which have increased
 anticoagulant activity and resistance to inactivation by serpins compared
 with the wild-type protein, while retaining the biological activity of
 the wild-type protein. A method of producing the derivatives using
 recombinant DNA methods is claimed. The protein C derivatives are useful
 for treating coronary syndromes and disease states predisposing to
 thrombosis (e.g., myocardial infarction and unstable angina), vascular
 occlusive disorders and hypercoagulable states, sepsis (in combination
 with bactericidal permeability increasing protein or with tissue factor
 pathway inhibitor), thrombotic disorders (in combination with an anti-
 platelet agent or by local delivery through an intracoronary catheter),
 protein C deficiency, acute arterial thrombotic occlusion,
 thromboembolism, or stenosis in coronary, cerebral or peripheral arteries
 or in vascular grafts. Human patients with genetically predisposed
 or thrombotic disorders may be treated by gene therapy (all claimed)

Sequence 419 AA;
 Query Match 88.9%; Score 176; DB 4; Length 419;
 Best Local Similarity 77.3%; Pred. No. 1.9e-20;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

1 ANSFLXLRHSSLRKXCIXICDPFXAKXIFEDVDPTLAFWSKH 44
 1 ANSFLXLRHSSLRKXCIXICDPFXAKXIFEDVDPTLAFWSKH 44

RESULT 3
 AAB82675
 ID AAB82675 standard; protein; 419 AA.
 AC AAB82675;

XX 15-OCT-2001 (first entry)
 XX XX Human protein C derivative (S1IG/G32E/N35D/L194S).
 XX XX Protein C; human; coronary syndrome; thrombosis; angina;
 XX myocardial infarction; vascular occlusive disorder; hypercoagulation;
 XX sepsis; protein C deficiency; occlusion; thromboembolism; stenosis;
 XX antibacterial; immunosuppressive; thrombolytic; cardiant; antiangiinal;
 XX anticoagulant; therapy; mutant; muten.
 XX Homo sapiens.
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 XX Domain 1..45
 XX /note= "Gla domain"
 XX Modified-site 6
 XX /note= "gamma-carboxylated"
 XX Modified-site 7
 XX /note= "gamma-carboxylated"
 XX Modified-site 11
 XX /note= "gamma-carboxylated"
 XX MISC-difference 14
 XX /note= "Ser in wild-type protein"
 XX Modified-site 16
 XX /note= "gamma-carboxylated"
 XX Modified-site 19
 XX /note= "gamma-carboxylated"
 XX Modified-site 20
 XX /note= "gamma-carboxylated"
 XX Modified-site 25
 XX /note= "gamma-carboxylated"
 XX Modified-site 26
 XX /note= "gamma-carboxylated"
 XX Modified-site 29
 XX /note= "gamma-carboxylated"
 XX MISC-difference 32
 XX /note= "N-glycosylated"
 XX MISC-difference 33
 XX /note= "Gln in wild-type protein"
 XX MISC-difference 33
 XX /note= "Asn in wild-type protein"
 XX Disulfide-bond 50..69
 XX /note= "Disulfide-bond 59..64"
 XX Disulfide-bond 80..89
 XX /note= "Disulfide-bond 98..109"
 XX Disulfide-bond 120..133
 XX /note= "Disulfide-bond 141..277"
 XX Disulfide-bond 156..157
 XX /note= "cleavage makes a 2-chain inactive precursor (155-
 XX amino acid light chain attached via a disulfide bond to a
 XX 262-amino acid heavy chain)"
 XX Peptide 158..169
 XX /note= "activation peptide; removal activates the 2-chain
 XX zymogen"
 XX Cleavage-site 169..170
 XX /note= "thrombin cleavage site"
 XX MISC-difference 194
 XX /note= "Leu in wild-type protein"
 XX Disulfide-bond 196..212
 XX /note= "N-glycosylated"
 XX Modified-site 313
 XX /note= "N-glycosylated"
 XX Modified-site 329
 XX /note= "N-glycosylated"
 XX Disulfide-bond 331..345
 XX Disulfide-bond 356..384
 XX W0200157193-A2.
 XX 09-AUG-2001.
 XX 19-JAN-2001; 2001MO-US000020.

XX 02-FEB-2000; 2000US-0179901P.
 PR 14-MAR-2000; 2000US-0189197B.
 XX (EHL) LILLY & CO EHL.
 XX Gerlitz BE, Jones BE,
 XX WPI; 2001-496919/54.
 DR N-PSDB; AAH26363.
 XX Novel human protein C derivative for treating, e.g., myocardial
 PT infarction, unstable angina, sepsis, thrombotic disorders, acute arterial
 PT thrombotic occlusion, and thromboembolism.
 XX Claim 3; Page 52-53; 63pp; English.
 XX The present sequence is that of a claimed human protein C derivative in
 CC which Ser at amino acid position 11 of the mature wild-type protein C
 CC sequence (see AAB82673) is substituted with Gly, Gln at position 33 with
 CC Gln, Asn at position 33 with Asp, and Leu at position 194 with Ser. The
 CC protein is an example of protein C derivatives of the invention that have
 CC at least 2 amino acid substitutions, but which have increased
 CC anticoagulant activity and resistance to inactivation by serpins compared
 CC with the wild-type protein, while retaining the biological activity of
 CC the wild-type protein. A method of producing the derivatives using
 CC recombinant DNA methods is claimed. The protein C derivatives are useful
 CC for treating coronary syndromes and disease states predisposing to
 CC thrombosis (e.g., myocardial infarction and unstable angina), vascular
 CC occlusive disorders and hypercoagulable states, sepsis (in combination
 CC with bactericidal permeability increasing protein or with tissue factor
 CC pathway inhibitor), thrombotic disorders (in combination with an anti-
 CC platelet agent or by local delivery through an intracoronary catheter),
 CC protein C deficiency, acute arterial thrombotic occlusion,
 CC thromboembolism, or stenosis in coronary, cerebral or peripheral arteries
 CC or in vascular grafts. Human patients with genetically predisposed
 CC prothrombotic disorders may be treated by gene therapy (all claimed)
 XX
 SQ Sequence 419 AA;
 Query Match 88.9%; Score 176; DB 4; Length 419;
 Best Local Similarity 77.3%; Pred. No. 1,9e-20;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 QY 1 ANSEFLXLRHSSILKRCYICDFFXXAKXIFEDVDITAFMSKH 44
 Db 1 ANSEFLXLRHSSILKRCYICDFFXXAKXIFEDVDITAFMSKH 44
 ID AAE08628 standard; protein; 419 AA.
 AC AAE08628;
 XX 01-NOV-2001 (first entry)
 XX Human protein C derivative #2.
 XX Human; protein C derivative; anticoagulation activity; thrombosis;
 XX serpin inactivation; acute coronary syndrome; myocardial infarction;
 XX vascular occlusive disorder; hypercoagulable state; angina; sepsis;
 XX disseminated intravascular coagulation; DIC; burn; transplantation;
 XX sickle cell disease; viral haemorrhagic fever; protein C deficiency;
 XX haemolytic uremic syndrome; acute arterial thrombotic occlusion;
 XX thromboembolism; prothrombotic disorder; gene therapy; thalassemia.
 XX Homo sapiens.
 OS Homo sapiens.
 XX W0200159084-A1.
 XX 16-AUG-2001.

PF 02-FEB-2001; 2001MO-US001221.
 XX 11-FEB-2000; 2000US-0181948P.
 PR 14-MAR-2000; 2000US-0189159P.
 XX (ELIL) LILLY & CO ELI.
 PA Gerlitz BE, Grinnell BW, Jones BE;
 PI WPI; 2001-514662/56.
 DR N-PSDB; AAD15225.
 XX
 PT Protein C derivative for treating acute coronary syndromes, vascular
 PT occlusive disorders, thrombotic disorders and sepsis, comprises
 PT substitutions at specified amino acid positions.
 XX
 PS Claim 4; Page 47-48; 59pp; English.
 XX
 CC The invention relates to human protein C derivatives and nucleic acid
 CC molecules encoding such derivatives. These derivatives have increased
 CC anticoagulation activity, resistance to sepsin inactivation and increased
 CC sensitivity to thrombin activation compared to wild type protein C, and
 CC retains the biological activity of the wild type human protein C. Protein
 CC C derivatives are useful in the manufacture of a medicament for the
 CC treatment of acute coronary syndromes e.g. myocardial infarction and
 CC unstable angina; and disease states predisposing to thrombosis; vascular
 CC occlusive disorders and hypercoagulable states e.g. disseminated
 CC intravascular coagulation (DIC), burns, transplantations, thalassaemia,
 CC sickle cell disease, viral haemorrhagic fever and haemolytic uremic
 CC syndrome; sepsis in combination with bacterial permeability increasing
 CC protein; thrombotic disorders in combination with an anti-platelet agent;
 CC protein C deficiency; acute arterial thrombotic occlusion,
 CC thromboembolism or stenosis in coronary, cerebral or peripheral arteries
 CC or in vascular grafts in combination with a thrombolytic agent. Nucleic
 CC acid molecules of the invention are useful for treating humans with
 CC genetically predisposed prothrombotic disorders by gene therapy. The
 CC present sequence is human protein C derivative
 SQ Sequence 419 AA;
 Query Match 88.9%; Score 176; DB 4; Length 419;
 Best Local Similarity 77.3%; Pred. No. 1.9e-20;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 QY 1 ANSFLXLRHSSLRXCIXXICDPFXAXXIFEDVDPTLAFWSKH 44
 DB 1 ANSFLXLRHSSLRXCIXXICDPFXAXXIFEDVDPTLAFWSKH 44
 RESULT 5
 AAE08627 ID AAE08627 standard; protein; 419 AA.
 XX
 AC AAE08627;
 DT 01-NOV-2001 (first entry)
 XX
 DE Human protein C derivative #1.
 XX
 KW Human; protein C derivative; anticoagulation activity; thrombosis;
 KW sepsin inactivation; acute coronary syndrome; myocardial infarction;
 KW vascular occlusive disorder; hypercoagulable state; angina; sepsis;
 KW disseminated intravascular coagulation; DIC; burn; transplantation;
 KW sickle cell disease; viral haemorrhagic fever; protein C deficiency;
 KW haemolytic uremic syndrome; acute arterial thrombotic occlusion;
 KW thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.
 OS Homo sapiens.
 XX
 PN WO200159084-A1.
 XX
 PD 16-AUG-2001.
 XX

PF 02-FEB-2001; 2001MO-US001221.
 XX 11-FEB-2000; 2000US-0181948P.
 PR 14-MAR-2000; 2000US-0189159P.
 XX (ELIL) LILLY & CO ELI.
 PA Gerlitz BE, Grinnell BW, Jones BE;
 PI WPI; 2001-514662/56.
 DR N-PSDB; AAD15225.
 XX
 PT Protein C derivative for treating acute coronary syndromes, vascular
 PT occlusive disorders, thrombotic disorders and sepsis, comprises
 PT substitutions at specified amino acid positions.
 XX
 PS Claim 3; Page 46-47; 59pp; English.
 XX
 CC The invention relates to human protein C derivatives and nucleic acid
 CC molecules encoding such derivatives. These derivatives have increased
 CC anticoagulation activity, resistance to sepsin inactivation and increased
 CC sensitivity to thrombin activation compared to wild type protein C, and
 CC retains the biological activity of the wild type human protein C. Protein
 CC C derivatives are useful in the manufacture of a medicament for the
 CC treatment of acute coronary syndromes e.g. myocardial infarction and
 CC unstable angina; and disease states predisposing to thrombosis; vascular
 CC occlusive disorders and hypercoagulable states e.g. disseminated
 CC intravascular coagulation (DIC), burns, transplantations, thalassaemia,
 CC sickle cell disease, viral haemorrhagic fever and haemolytic uremic
 CC syndrome; sepsis in combination with bacterial permeability increasing
 CC protein; thrombotic disorders in combination with an anti-platelet agent;
 CC protein C deficiency; acute arterial thrombotic occlusion,
 CC thromboembolism or stenosis in coronary, cerebral or peripheral arteries
 CC or in vascular grafts in combination with a thrombolytic agent. Nucleic
 CC acid molecules of the invention are useful for treating humans with
 CC genetically predisposed prothrombotic disorders by gene therapy. The
 CC present sequence is human protein C derivative
 SQ Sequence 419 AA;
 Query Match 88.9%; Score 176; DB 4; Length 419;
 Best Local Similarity 77.3%; Pred. No. 1.9e-20;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 QY 1 ANSFLXLRHSSLRXCIXXICDPFXAXXIFEDVDPTLAFWSKH 44
 DB 1 ANSFLXLRHSSLRXCIXXICDPFXAXXIFEDVDPTLAFWSKH 44
 RESULT 6
 AAE08629 ID AAE08629 standard; protein; 419 AA.
 XX
 AC AAE08629;
 DT 01-NOV-2001 (first entry)
 XX
 DE Human protein C derivative #3.
 XX
 KW Human; protein C derivative; anticoagulation activity; thrombosis;
 KW sepsin inactivation; acute coronary syndrome; myocardial infarction;
 KW vascular occlusive disorder; hypercoagulable state; angina; sepsis;
 KW disseminated intravascular coagulation; DIC; burn; transplantation;
 KW sickle cell disease; viral haemorrhagic fever; protein C deficiency;
 KW haemolytic uremic syndrome; acute arterial thrombotic occlusion;
 KW thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.
 OS Homo sapiens.
 XX
 PN Key Location/Qualifiers
 FT Misc-difference 10
 FT /note= "Encoded by CAA"
 FT
 XX

PN W0200159084-A1.
 XX 16-AUG-2001.
 XX 02-FEB-2001; 2001MO-US001221.
 XX 11-FEB-2000; 2000US-0181948P.
 PR 14-MAR-2000; 2000US-0189199P.
 XX (ELIL) LILLY & CO ELI.
 PA Geriltz BE, Grinnell BW, Jones BE;
 XX WPI; 2001-514662/56.
 DR N-PSDB; AADI5227.
 XX Protein C derivative for treating acute coronary syndromes, vascular
 PT occlusive disorders, thrombotic disorders and sepsis, comprises
 PT substitutions at specified amino acid positions.
 XX Claim 5; Page 48-49; 59pp; English.
 CC The invention relates to human protein C derivatives and nucleic acid
 CC molecules encoding such derivatives. These derivatives have increased
 CC anticoagulation activity, resistance to serpin inactivation and increased
 CC sensitivity to thrombin activation compared to wild type protein C, and
 CC retains the biological activity of the wild type human protein C. Protein
 CC derivatives are useful in the manufacture of a medicament for the
 CC treatment of acute coronary syndromes e.g. myocardial infarction and
 CC unstable angina; and disease states predisposing to thrombosis; vascular
 CC occlusive disorders and hypercoagulable states e.g. disseminated
 CC intravascular coagulation (DIC), burns, transplantations, thalassaemia,
 CC sickle cell disease, viral haemorrhagic fever and haemolytic uremic
 CC syndrome; sepsis in combination with bacterial permeability increasing
 CC protein; thrombotic disorders in combination with an anti-platelet agent;
 CC thromboembolism or stenosis in coronary, cerebral or peripheral arteries
 CC or in vascular grafts in combination with a thrombolytic agent. Nucleic
 CC acid molecules of the invention are useful for creating humans with
 CC genetically predisposed prothrombotic disorders by gene therapy. The
 CC present sequence is human protein C derivative
 SQ Sequence 419 AA;
 QY Query Match 88.9%; Score 176; DB 4; Length 419;
 DB Best Local Similarity 77.3%; Pred. No. 1.9e-20;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 1 ANSFLXXLRHSSLSRXICIXXICDFFXXAXXIFEDVDTTAFWSKH 44
 1 ANSFLLELRHGSLEKCEICICDFEAKXIFEDVDTTAFWSKH 44
 RESULT 7
 ID ABB79950 standard; protein; 45 AA.
 XX ABB79950;
 XX 12-DEC-2002 (first entry)
 DB Human protein C mutated Gla domain SED.
 XX Protein C; Gla domain; human; blood clotting; anticoagulant;
 KM thrombolytic; antiarteriosclerotic; cardiant; antiagregant; mutant;
 XX mutein.
 XX Homo sapiens.
 OS Synthetic.
 XX Key Location/Qualifiers
 FH Msc-difference 23 /note= "wild-type Asp substituted by Ser"
 FT

FT Msc-difference 32 /note= "wild-type Gln substituted by Glu"
 FT Msc-difference 33 /note= "wild-type Asn substituted by Asp"
 FT /note= "wild-type Asp substituted by Asp"
 XX W0200270681-A1.
 XX 12-SEP-2002.
 XX 01-MAR-2002; 2002MO-SE000363.
 XX 02-MAR-2001; 2001US-0272466P.
 XX (TACT-) TAC THROMBOSIS & COAGULATION AB.
 PA Dahlback B;
 XX WPI; 2002-713449/77.
 DR New variant blood coagulation component, useful for manufacturing a
 PT medicament for treating or preventing coagulation disorders, e.g.
 PT thrombosis, comprises an anticoagulant activity in the protein C-
 PT anticoagulant system of blood.
 XX Example 1; Page; 56pp; English.
 PS The present sequence is the protein sequence of a mutated Gla domain (N-
 CC terminal amino acids 1-45) of human protein C. The mutated Gla domain
 CC contains the substitution mutations D23S, Q32E and N33D. Protein C and
 CC activated protein C variants comprising a mutated Gla domain are provided
 CC by the invention. The variants contain at least 6, and optionally 7-10,
 CC amino acid substitutions. A preferred mutant (designated QGNS2D, see
 CC ABB79946) has the mutations H10Q, S19G, S12N, D23S, Q32E, N33D and H44Y,
 CC and shows greatly enhanced anticoagulant activity in standard in vitro
 CC coagulation assays. The present mutant (designated SED) was produced in
 CC an example from the invention as a step toward the production of the
 CC QGNS2D mutant Gla domain, and shows little, if any, improvement in
 CC anticoagulant activity over wild-type activated protein C. The invention
 CC provides methods for producing the variants based on DNA technology, and
 CC with the use of the variants for the treatment of coagulation disorders
 CC such as thrombosis or Apc resistance, or in diagnostic test systems for
 CC assaying components of the protein C-anticoagulant system (all claimed).
 CC The variants may also be used in treating arteriosclerosis, myocardial
 CC infarction, and disseminated intravascular coagulation. Note: The present
 CC sequence is not shown in the specification but is derived from the human
 CC wild-type Gla domain sequence given on page 7 of the specification (see
 CC ABB79947)
 SQ Sequence 45 AA;
 QY Query Match 87.9%; Score 174; DB 5; Length 45;
 DB Best Local Similarity 77.3%; Pred. No. 3.6e-21;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 1 ANSFLXXLRHSSLSRXICIXXICDFFXXAXXIFEDVDTTAFWSKH 44
 1 ANSFLLELRHGSLEKCEICICDFEAKXIFEDVDTTAFWSKH 44
 RESULT 8
 ID AAY18299 standard; peptide; 44 AA.
 XX AAY18299;
 XX 17-AUG-1999 (first entry)
 DB Modified GLA domain of vitamin K-dependent protein.
 XX Gla domain; mutein; vitamin K-dependent protein; clotting disorder;
 KM therapy.
 XX Homo sapiens.
 OS

OS Synthetic.
 XX Key Location/Qualifiers
 FT Misc-difference 1..44
 FT /note= "Xaa= gamma-carboxyglutamic acid, or glutamic
 FT acid"
 XX
 XX
 PN WO9920767-A1.
 XX
 XX 29-APR-1999.
 PD
 XX 20-OCT-1998; 98WO-US022152.
 PF
 XX 23-OCT-1997; 97US-00955636.
 PR
 XX (MINU) UNIV MINNESOTA.
 PA
 XX Nelastuen GL;
 PI
 XX WPI; 1999-288309/24.
 DR
 XX
 XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid
 PT domain, useful for treating clotting disorders.
 XX
 XX Claim 8; Page 78; 86pp; English.
 PS
 XX This sequence represents a modified GLA (gamma-carboxyglutamic acid)
 CC domain. The invention relates to a vitamin K-dependent polypeptide
 CC comprising a modified GLA domain containing an amino acid substitution
 CC which enhances membrane binding of the modified polypeptide as compared
 CC to the native polypeptide. The polypeptide is used to treat a clotting
 CC disorder by decreasing or increasing clot formation. Modification of the
 CC GLA domain results in a protein which has enhanced membrane binding
 CC affinity as compared to the native protein
 CC
 SQ Sequence 44 AA;
 Query Match 86.9%; Score 172; DB 2; Length 44;
 Best Local Similarity 97.7%; Pred. No. 7.6e-21;
 Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ANSFLXXLRHSLRXKXICIXXICDFXXAKXIFEDVDPTLAFWSKH 44
 DB 1 ANSFLXXLRHSLRXKXICIXXICDFXXAKXIFEDVDPTLAFWSKH 44
 RESULT 9
 ID AAY18309 standard; peptide; 44 AA.
 XX
 XX AAY18309;
 AC
 XX 17-AUG-1999 (first entry)
 DT
 XX Modified GLA domain of vitamin K-dependent protein.
 DE
 XX GLA domain; mutein; vitamin K-dependent protein; clotting disorder;
 KM therapy.
 KM
 XX Homo sapiens.
 OS
 XX Synthetic.
 OS
 XX Key Location/Qualifiers
 FT Misc-difference 1..44
 FT /note= "Xaa= gamma-carboxyglutamic acid, or glutamic
 FT acid"
 XX
 XX
 PN WO9920767-A1.
 XX
 XX 29-APR-1999.
 PD
 XX 20-OCT-1998; 98WO-US022152.
 PF
 XX

PR 23-OCT-1997; 97US-00955636.
 XX
 XX (MINU) UNIV MINNESOTA.
 PA
 XX Nelastuen GL;
 PI
 XX WPI; 1999-288309/24.
 DR
 XX
 XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid
 PT domain, useful for treating clotting disorders.
 XX
 XX Disclosure; Page 79-80; 86pp; English.
 PS
 XX This sequence represents a modified GLA (gamma-carboxyglutamic acid)
 CC domain. The invention relates to a vitamin K-dependent polypeptide
 CC comprising a modified GLA domain containing an amino acid substitution
 CC which enhances membrane binding of the modified polypeptide as compared
 CC to the native polypeptide. The polypeptide is used to treat a clotting
 CC disorder by decreasing or increasing clot formation. Modification of the
 CC GLA domain results in a protein which has enhanced membrane binding
 CC affinity as compared to the native protein
 CC
 SQ Sequence 44 AA;
 Query Match 86.9%; Score 172; DB 2; Length 44;
 Best Local Similarity 97.7%; Pred. No. 7.6e-21;
 Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 ANSFLXXLRHSLRXKXICIXXICDFXXAKXIFEDVDPTLAFWSKH 44
 DB 1 ANSFLXXLRHSLRXKXICIXXICDFXXAKXIFEDVDPTLAFWSKH 44
 RESULT 10
 ID AAY18298 standard; peptide; 44 AA.
 XX
 XX AAY18298;
 AC
 XX 17-AUG-1999 (first entry)
 DT
 XX Modified GLA domain of vitamin K-dependent protein.
 DE
 XX GLA domain; mutein; vitamin K-dependent protein; clotting disorder;
 KM therapy.
 KM
 XX Homo sapiens.
 OS
 XX Synthetic.
 OS
 XX Key Location/Qualifiers
 FT Misc-difference 1..44
 FT /note= "Xaa= gamma-carboxyglutamic acid, or glutamic
 FT acid"
 XX
 XX
 PN WO9920767-A1.
 XX
 XX 29-APR-1999.
 DT
 XX 20-OCT-1998; 98WO-US022152.
 PF
 XX 23-OCT-1997; 97US-00955636.
 PR
 XX (MINU) UNIV MINNESOTA.
 PA
 XX Nelastuen GL;
 PI
 XX WPI; 1999-288309/24.
 DR
 XX
 XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid
 PT domain, useful for treating clotting disorders.
 XX
 XX Claim 7; Page 78; 86pp; English.

CC This sequence represents a modified GLA (gamma-carboxyglutamic acid)
 CC domain. The invention relates to a vitamin K-dependent polypeptide
 CC comprising a modified GLA domain containing an amino acid substitution
 CC which enhances membrane binding of the modified polypeptide as compared
 CC to the native polypeptide. The polypeptide is used to treat a clotting
 CC disorder by decreasing or increasing clot formation. Modification of the
 CC GLA domain results in a protein which has enhanced membrane binding
 CC affinity as compared to the native protein
 CC
 XX
 SQ Sequence 44 AA;

Query Match 86.9%; Score 172; DB 2; Length 44;
 Best Local Similarity 97.7%; Pred. No. 7.6e-21;
 Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHSSLRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 1 ANSFLXXLRHSSLRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 Db

RESULT 11
 AAY18303
 ID AAY18303 standard; peptide; 44 AA.

XX AC AAY18303;

XX DT 17-APR-1999 (first entry)

XX DE Human protein C GLA domain.

XX KW GLA domain; vitamin K-dependent protein; clotting disorder; therapy.

XX OS Homo sapiens.

XX FT Key Location/Qualifiers

FT MISC-difference 1..44 /note="Xaa= gamma-carboxyglutamic acid, or glutamic
 FT acid"

XX PV WO9920767-A1.

XX PD 29-APR-1999.

XX PF 20-OCT-1998; 98WO-US022152.

XX PR 23-OCT-1997; 97US-00955636.

XX PA (MINU) UNIV MINNESOTA.

XX PI Nelsestuen GL;

XX DR WPI; 1999-288309/24.

XX PT Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid
 XX domain, useful for treating clotting disorders.

XX PS Disclosure; Page 14; 86pp; English.

XX This sequence is the protein C GLA (gamma-carboxyglutamic acid) domain.
 CC The invention relates to a vitamin K-dependent polypeptide comprising a
 CC modified GLA domain containing an amino acid substitution which enhances
 CC membrane binding of the modified polypeptide as compared to the native
 CC polypeptide. The polypeptide is used to treat a clotting disorder by
 CC decreasing or increasing clot formation. Modification of the GLA domain
 CC results in a protein which has enhanced membrane binding affinity as
 CC compared to the native protein
 CC
 XX

SQ Sequence 44 AA;

Query Match 86.9%; Score 172; DB 2; Length 44;
 Best Local Similarity 95.5%; Pred. No. 7.6e-21;
 Matches 42; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHSSLRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 1 ANSFLXXLRHSSLRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 Db

RESULT 12
 AAB36402
 ID AAB36402 standard; peptide; 44 AA.

XX AC AAB36402;

XX DT 27-FEB-2001 (first entry)

XX DE Human protein C gamma-carboxyglutamic acid domain SEQ ID NO:1.

XX KW Vitamin K-dependent protein; factor VII; protein C; GLA domain;
 KW gamma-carboxyglutamic acid domain; factor IX; protein S; protein Z;
 KW factor X; prothrombin; enhanced membrane binding affinity;
 KW clot formation; thrombolytic; haemostatic; bleeding disorder; thrombosis;
 KW clotting disorder; haemophilia A; haemophilia B; liver disease.

XX OS Homo sapiens.

XX PN WO200066753-A2.

XX PD 09-NOV-2000.

XX PF 28-APR-2000; 2000WO-US011416.

XX PR 29-APR-1999; 99US-00302239.

XX PA (MINU) UNIV MINNESOTA.

XX PI Nelsestuen GL;

XX DR WPI; 2001-007226/01.

XX PT Novel vitamin K-dependent polypeptide useful for treating clotting
 XX disorders such as thrombosis and hemophilia, comprises modified gamma-
 XX carboxy glutamic acid domain that enhances membrane binding affinity.

XX PS Example 5; Page 42; 81pp; English.

XX The present invention describes a vitamin K-dependent polypeptide (I)
 CC comprising a modified gamma-carboxy glutamic acid (GLA) domain having at
 CC least one amino acid substitution, that enhances membrane binding
 CC affinity and the activity of the polypeptide relative to a corresponding
 CC native vitamin K-dependent polypeptide and inhibits clot formation. (II)
 CC can have thrombolytic and haemostatic activities, and can be used as an
 CC inhibitor of clot formation. (II) is useful for decreasing clot formation
 CC in a mammal, a factor VII or factor IX containing a modified GLA domain
 CC is useful for increasing clot formation and for treating a bleeding
 CC disorder including thrombosis and clotting disorders such as haemophilia
 CC A, haemophilia B and liver disease. The present sequence represents a
 CC human protein C GLA domain sequence, given in the exemplification of the
 CC present invention
 CC
 XX

SQ Sequence 44 AA;

Query Match 86.9%; Score 172; DB 4; Length 44;
 Best Local Similarity 95.5%; Pred. No. 7.6e-21;
 Matches 42; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHSSLRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 1 ANSFLXXLRHSSLRXCIXXICDFXXAKXIFONVDTLAFWSKH 44
 Db

RESULT 13

ADD50094
 ID ADD50094 standard; protein; 44 AA.

XX AC ADD50094;

```

XX 15-JAN-2004 (first entry)
DT Human vitamin K-dependent protein #1.
DE
XX Human, vitamin K-dependent protein; gamma-carboxyglutamic acid domain;
XX GLA domain; membrane binding affinity; clot formation; haemophilia;
XX clotting disorder; site directed mutagenesis; haemostatic; anticoagulant;
XX thrombolytic.
XX
XX Homo sapiens.
OS
XX US2003100506-A1.
XX
XX 29-MAY-2003.
XX
XX 18-NOV-2002; 2002US-00298330.
XX
XX 23-OCT-1997; 97US-00955636.
XX
XX 29-APR-1998; 98US-00302239.
XX
XX 03-FEB-2000; 2000US-00497591.
XX
XX (NEUS/) NEUSESTUEN G L.
XX
XX Nelstuen GL;
XX
XX WPI; 2003-606646/57.
XX
XX New vitamin K-dependent polypeptide for modulating clot formation in
XX mammals comprises a modified gamma-carboxyglutamic acid domain that
XX enhances membrane binding affinity and activity of the polypeptide.
XX
XX Example 5; SEQ ID NO 1; 51pp; English.
XX
XX The invention relates to a vitamin K-dependent polypeptide comprising a
XX modified gamma-carboxyglutamic acid (GLA) domain that enhances membrane
XX binding affinity and activity of the polypeptide relative to a
XX corresponding native vitamin K-dependent polypeptide, where the modified
XX GLA domain comprises a glutamic acid residue at position 34. The
XX polypeptide is useful in modulating clot formation in mammals or in
XX treating certain types of haemophilia or clotting disorders. The membrane
XX binding affinity of polypeptides is increased by site directed
XX mutagenesis in the GLA domain. This sequence represents a vitamin K-
XX dependent protein of the invention.
XX
XX Sequence 44 AA;
SQ
XX
XX Query Match 86.9%; Score 172; DB 7; Length 44;
XX Best Local Similarity 95.5%; Pred. No. 7.6e-21;
XX Matches 42; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 ANSFLXLRHSSILRXICIXICDPFXAKXIFEDVDTLAFWSKH 44
XX 1 ANSFLXLRHSSILRXICIXICDPFXAKXIFQVNDTLAFWSKH 44
XX
XX RESULT 14
XX AAW75710
XX ID AAW75710 standard; protein; 45 AA.
XX
XX AAW75710;
XX
XX 08-DEC-1998 (first entry)
XX
XX Partial human protein C amino acid sequence.
XX
XX Gamma carboxyglutamic acid; human protein C; GLA domain; chimera;
XX PRC/RSV; RSV-PC; amplification; PCR; primer; transfection; anticoagulant;
XX human 293 cell; Protein S; myocardial infarction; venous thrombosis;
XX disseminated intravascular coagulation; thromboembolic disease; lupus;
XX adult respiratory distress syndrome; factor V Leiden; stroke.
XX
XX Homo sapiens.
OS

```

```

XX Key Location/Qualifiers
XX Misc-difference 6 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 7 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 14 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 16 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 19 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 20 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 25 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 26 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 29 /note= "Gamma carboxyglutamic acid"
XX Misc-difference 29 /note= "Gamma carboxyglutamic acid"
XX
XX MO982018-A1.
XX
XX 14-MAY-1998.
XX
XX 07-NOV-1997; 97WO-US020376.
XX
XX 08-NOV-1996; 96US-00745254.
XX
XX 25-JUL-1997; 97US-0053768P.
XX
XX (OKLA-) OKLAHOMA MEDICAL RES FOUND.
XX
XX Bsmen CT, Smitrov M;
XX
XX WPI; 1998-286934/25.
XX
XX Protein C chimeric proteins for use as anticoagulants - having gamma
XX carboxyglutamic acid region replaced with Vitamin K dependent clotting
XX factor e.g. prothrombin.
XX
XX Example 1; Page 15; 42pp; English.
XX
XX The present sequence represents the first three exons of the human
XX protein C protein, which contains gamma carboxyglutamic acid modified
XX residues. This sequence was replaced with the corresponding regions of
XX modified human prothrombin (AAW75709), to create a protein C prothrombin
XX GLA domain chimera. To produce this chimera, the wild-type protein C cDNA
XX was ligated into pRC/RSV to form RSV-PC, and then was digested with
XX restriction enzymes to remove the first three exons and the first codon
XX of exon four. The prothrombin cDNA was amplified, digested, and then
XX ligated into RSV-PC at the identical site where the protein C cDNA exons
XX 1-3 had been removed. This construct was then transfected into human 293
XX cells, from which the chimeric protein can be purified. This chimeric
XX protein can be used as an anticoagulant, to treat disorders where protein
XX S is low, some forms of lupus, following stroke or myocardial infarction,
XX after venous thrombosis and in disseminated intravascular coagulation,
XX adult respiratory distress syndrome, in thromboembolic disease or factor
XX V Leiden
XX
XX Sequence 45 AA;
SQ
XX
XX Query Match 86.9%; Score 172; DB 2; Length 45;
XX Best Local Similarity 95.5%; Pred. No. 7.8e-21;
XX Matches 42; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
XX
XX 1 ANSFLXLRHSSILRXICIXICDPFXAKXIFEDVDTLAFWSKH 44
XX 1 ANSFLXLRHSSILRXICIXICDPFXAKXIFQVNDTLAFWSKH 44
XX
XX RESULT 15
XX ABB79947
XX ID ABB79947 standard; protein; 45 AA.
XX

```

```

XX      ABB79947;
AC
XX
XX      12-DEC-2002 (first entry)
DT
XX
XX      Human protein C Gla domain.
DE
XX
XX      Protein C; Gla domain; human; blood clotting; anticoagulant;
KW      thrombolytic; antiarteriosclerotic; cardiant; antiagregant.
XX
XX      Homo sapiens.
OS
XX      MO200270681-A1.
PN
XX      12-SEP-2002.
PD
XX      01-MAR-2002; 2002WC-SE000363.
PF
XX      02-MAR-2001; 2001US-027246P.
PR
XX
XX      (TACT-) TAC THROMBOSIS & COAGULATION AB.
PA
XX      Dahlback B;
PI
XX
XX      WPI; 2002-713449/77.
DR
XX
XX      New variant blood coagulation component, useful for manufacturing a
PT      medicament for treating or preventing coagulation disorders, e.g.
PT      thrombosis, comprises an anticoagulant activity in the protein C-
PT      anticoagulant system of blood.
XX
XX      Disclosure; Page 7; 58p; English.
PS
XX
XX      The present sequence is the protein sequence of the Gla domain (N-
CC      terminal amino acids 1-45) of human protein C. The invention provides
CC      human protein C and activated protein C variants in which the Gla domain
CC      contains at least 6, and optionally 7-10, amino acid substitutions
CC      preferably the substitution mutations H10Q, S11G, S12N, D35E, Q32E, N33D
CC      and H44Y (see ABB79946). Protein C variants comprising the mutated Gla
CC      domain show much enhanced anticoagulant activity, as shown in increased
CC      clotting time in standard in vitro coagulation assays, as a result of
CC      enhanced calcium and/or membrane binding properties. The invention
CC      provides methods for producing the variants based on DNA technology, and
CC      for using the variants in the treatment of coagulation disorders such as
CC      thrombosis or APC resistance, or in diagnostic test systems for assaying
CC      components of the protein C-anticoagulant system (all claimed). The
CC      variants may also be used in treating arteriosclerosis, myocardial
CC      infarction, and disseminated intravascular coagulation
XX
XX      Sequence 45 AA;
SQ

```

```

Query Match      86.9%; Score 172; DB 5; Length 45;
Best Local Similarity 75.0%; Pred. No. 7, 8e-21;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

```

```

QY      1 ANSFLXXLRSSSLRXCIXXCDFXXKXIFEDVDDTLAFWSKH 44
      |||||
Db      1 ANSFLSELRRSSSLRXCIEICDFEFAKXIFQVDDTLAFWSKH 44

```

Search completed: March 1, 2004, 10:01:23
Job time : 52 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 1, 2004, 09:59:33 ; Search time 13.5 Seconds
(without alignments)
313.513 Million cell updates/sec

Title: SEQ1-32GLU-33ASP

Perfect score: 198

Sequence: 1 ANSFLXLRHSLXRCIXX.....XXAKXIFedVDDTLAFWSKH 44

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues

Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: PIR 78:**

1: PIR1:**

2: PIR2:**

3: PIR3:**

4: PIR4:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	172	86.9	461	1 KXHU	protein C (activat
2	133	67.2	461	1 JX0210	protein C (activat
3	132	66.7	461	1 S18994	protein C (activat
4	115	58.1	456	1 KXBO	protein C (activat
5	107	54.0	482	1 EXRT	coagulation factor
6	103	52.0	492	1 EXBO	coagulation factor
7	102	51.5	488	1 EXHU	coagulation factor
8	94	47.5	443	2 I46932	coagulation factor
9	92	46.5	466	1 KFHU7	coagulation factor
10	78.5	39.6	617	2 S10511	thrombin (EC 3.4.2
11	78.5	39.6	618	2 A35827	thrombin (EC 3.4.2
12	78	39.4	407	1 KFB07	coagulation factor
13	76	38.4	622	1 TBHU	thrombin (EC 3.4.2
14	75	37.9	475	1 EXCH	coagulation factor
15	74	37.4	642	2 S33434	plasma protein S p
16	74	37.4	676	1 KXHU5	plasma protein S p
17	73	36.9	452	1 A30351	coagulation factor
18	73	36.9	459	2 J00419	coagulation factor
19	73	36.9	646	2 S38819	plasma protein S -
20	72	36.4	675	1 KXBO5	plasma protein S p
21	70	35.4	675	1 KXHU5	plasma protein S p
22	69	34.8	461	1 KFHU7	coagulation factor
23	67	33.8	642	2 S33433	plasma protein S p
24	66	33.3	416	1 KFB0	coagulation factor
25	64	32.3	625	1 TBBO	thrombin (EC 3.4.2
26	63	31.8	675	1 KXWSS	plasma protein S p
27	61.5	31.1	396	1 KXBOZ	plasma protein Z -
28	57.5	29.0	422	1 KXHUZ	plasma protein Z p
29	56.5	28.5	594	2 D84859	probable MAP kinas

RESULT 1

KXHU

protein C (activated) (EC 3.4.21.69) precursor - human

N;Alternate names: autoprothrombin IIA; plasma protein C

C;Species: Homo sapiens (man)

C;Date: 17-Mar-1987 #sequence revision 17-Mar-1987 #text_change 16-Jul-1999

C;Accession: A22331; A25426; A21781; A23789; A00927

R;Poster, D.C.; Yoshitake, S.; Davie, E.W.

Proc. Natl. Acad. Sci. U.S.A. 82, 4673-4677, 1985

A;Title: The nucleotide sequence of the gene for human protein C.

A;Reference number: A22331; MUID:85270390; PMID:2991887

A;Accession: A22331

A;Molecule type: DNA

A;Residues: 1-461 <POS1>

A;Cross-references: GB:M11228; NID:gl90333; PIDN:AAA60166.1; PID:gl90334

R;Plutsky, J.; Hoskins, J.A.; Long, G.L.; Crabtree, G.R.

Proc. Natl. Acad. Sci. U.S.A. 83, 546-550, 1986

A;Title: Evolution and organization of the human protein C gene.

A;Reference number: A25426; MUID:86120978; PMID:3511471

A;Accession: A25426

A;Molecule type: DNA

A;Residues: 1-445, 'L', 446-461 <PLU>

A;Cross-references: GB:M12712; NID:gl90330; PIDN:AAA60165.1; PID:gl90332

R;Poster, D.; Davie, E.W.

Proc. Natl. Acad. Sci. U.S.A. 81, 4766-4770, 1984

A;Title: Characterization of a cDNA coding for human protein C.

A;Reference number: A21781; MUID:84272714; PMID:6589623

A;Accession: A21781

A;Molecule type: mRNA

A;Residues: 'Q', 107-461 <POS2>

A;Cross-references: GB:X02059; NID:gl90322; PIDN:AAA60164.1; PID:gl90323

R;Beckmann, R.J.; Schmidt, R.J.; Sautter, R.F.; Plutsky, J.; Crabtree, G.R.; Long, G.L.

Nucleic Acids Res. 13, 5233-5247, 1985

A;Title: The structure and evolution of a 461 amino acid human protein C precursor and it

A;Reference number: A23789; MUID:85269639; PMID:2991859

A;Accession: A23789

A;Molecule type: mRNA

A;Residues: 1-461 <BEC>

A;Cross-references: GB:X02750; NID:G35689; PIDN:CAA26528.1; PID:g763120

R;Mileich, J.P.; Broze Jr., G.J.

J. Biol. Chem. 265, 11397-11404, 1990

A;Title: Beta protein C is not glycosylated at asparagine 329. The rate of translation me

A;Reference number: A4605; MUID:90293094; PMID:1694179

A;Contents: annotation; carbohydrate binding sites; activation peptide

A;Note: the alpha form of protein C is glycosylated at Asn-329, and the beta form is not

R;Harris, R.J.; Ling, V.T.; Spellman, M.W.

J. Biol. Chem. 267, 5102-5107, 1992

A;Title: O-linked fucose is present in the first epidermal growth factor domain of factor

A;Reference number: A4606; MUID:92184750; PMID:1544894

A;Contents: annotation; Beta-hydroxyaspartic acid

C;Comment: Protein C is the zymogen of the vitamin K-dependent serine proteinase that in

activation of factor Va is strongly enhanced by complexing with protein S. Protein C also fa

probable MAP kinas
probable MAP kinas
growth arrest-spec
growth potentialin
primosomal replica
growth arrest-spec
protein-tyrosine k
protein-tyrosine k
hypothetical prote
tyrosine kinase re
endothelial kinase
protein-tyrosine k
protein-tyrosine k
vascular endotheli
hypothetical prote
ammonium transport

C;Comment: Protein C is synthesized in the liver as a single chain precursor, which is dimerized, which cleaves a dodecapeptide from the amino end of the heavy chain; this reaction, C;Genetics: A;Accession: GDB:PROC A;Cross-references: GDB:120317; OMIM:176860 A;Map position: 2q33-q32 A;Introns: 24/1; 79/3; 88/1; 134/1; 179/1; 226/3; 266/1 C;Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology C;Keywords: anticoagulant; beta-hydroxyaspartic acid; blood coagulation; calcium binding F;1-32/Domain: signal sequence #status predicted <SIG> F;27-86/Domain: Gla domain homology <Gla> F;33-42/Domain: propeptide #status predicted <PRO> F;43-197/Domain: protein C light chain #status predicted <LCH> F;92-113/Domain: EGF homology <EG1> F;140-175/Domain: EGF homology <EG2> F;200-461/Domain: protein C heavy chain #status predicted <HCH> F;200-211/Domain: activation peptide #status experimental <APT> F;212-445/Domain: trypsin homology <TRY> F;48-49-56-61-62-67-68-71/Modified site: gamma-carboxyglutamic acid (Glu) #status exp F;59-64-92-105-120-122-131-140-151-147-160-162-175-183-319-238-254-373-387-398-426/D F;106-111/Disulfide bonds: #status predicted F;110/Binding site: carbohydrate (Thr) (covalent) #status absent F;113/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status experimental F;139-290-355/Binding site: carbohydrate (Asn) (covalent) #status experimental F;211-212/Cleavage site: Arg-Leu (thrombin) #status predicted F;253-299-402/Active site: His, Asp, Ser #status predicted F;371/Binding site: carbohydrate (Asn) (covalent) (partial) #status atypical

Query Match 86.9%; Score 172; DB 1; Length 461;
Best Local Similarity 75.0%; Pred. No. 1e-19;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLLXLRHSLRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44
DB 43 ANSFLEELRHSLEECIEECIECFEAEQIFQNVDDTLAFWSKH 96

RESULT 2
JX0210
Protein C (activated) (EC 3.4.21.69) precursor - mouse
N;Alternate names: vitamin K-dependent serine proteinase
C;Species: Mus musculus (house mouse)
C;Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 16-Jun-2000
C;Accession: JX0210
J;Tada, N.; Sato, M.; Tsujimura, A.; Iwase, R.; Hashimoto-Gotoh, T.
J. Biochem. 111, 491-495, 1992
A;Title: Isolation and characterization of a mouse protein C cDNA.
A;Reference number: JX0210; MUID:92316897; PMID:1618739
A;Accession: JX0210
A;Molecule type: mRNA
A;Residues: 1-461 <OK>
A;Cross-references: GB:D10445; NID:9220385; PIDN:BAA01235.1; PID:G220386
A;Experimental source: liver
C;Comment: Protein C is the zymogen of the vitamin K-dependent serine proteinase that re

C;Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
F;1-33/Domain: signal sequence #status predicted <SIG>
F;27-85/Domain: Gla domain homology <Gla>
F;34-41/Domain: propeptide #status predicted <PRO>
F;42-196-199-461/Domain: protein C #status predicted <PRC>
F;42-196/Domain: light chain #status predicted <PCL>
F;91-130/Domain: EGF homology <EG1>
F;139-174/Domain: EGF homology <EG2>
F;199-461/Domain: heavy chain #status predicted <ACT>
F;199-211/Domain: activation peptide #status predicted <ACT>
F;212-461/Domain: protein C #status predicted <PRC>
F;212-445/Domain: trypsin homology <TRY>
F;47-48-55-56-57-60-61-66-67-70-76/Modified site: gamma-carboxyglutamic acid (Glu) #status
F;112/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status predicted
F;121-130-139-150-146-159-161-174-182-319-238-254-373-387-398-426/Disulfide bonds: #stat
F;214-290-355/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;253-299-402/Active site: His, Asp, Ser #status predicted

Query Match 67.2%; Score 133; DB 1; Length 461;
Best Local Similarity 56.8%; Pred. No. 2e-13;
Matches 25; Conservative 7; Mismatches 12; Indels 0; Gaps 0;

QY 1 ANSFLLXLRHSLRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44
DB 42 ANSFLEELRHSLEECIEECIECFEAEQIFQNVDDTLAFWSKH 85

RESULT 3

S18994

Protein C (activated) (EC 3.4.21.69) precursor - rat

C;Species: Rattus norvegicus (Norway rat)

C;Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 29-Oct-1999

C;Accession: S18994; S24312

R;Okafuji, T.; Maekawa, K.; Nawa, K.; Marumoto, Y.

submitted to the EMBL Data Library, February 1992

A;Description: The cDNA cloning and mRNA expression of rat protein C.

A;Reference number: S18994

A;Accession: S18994

A;Status: preliminary

A;Molecule type: mRNA

A;Residues: 1-461 <OK>

R;Okafuji, T.; Maekawa, K.; Nawa, K.; Marumoto, Y.

Biochim. Biophys. Acta 1131, 329-332, 1992

A;Title: The cDNA cloning and mRNA expression of rat protein C.

A;Reference number: S24312; MUID:92329550; PMID:1627650

A;Accession: S24312

A;Status: preliminary

A;Molecule type: mRNA

A;Residues: 1-461 <OK>

C;Cross-references: EMBL:X64336; NID:956962; PIDN:CAA45617.1; PID:G56963

C;Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology

C;Keywords: beta-hydroxyaspartic acid; glycoprotein; hydrolase; serine proteinase

F;1-32/Domain: signal sequence #status predicted <SIG>

F;27-85/Domain: Gla domain homology <Gla>

F;33-42/Domain: propeptide #status predicted <PRO>

F;43-461/Domain: protein C #status predicted <PRC>

F;91-130/Domain: EGF homology <EG1>

F;139-174/Domain: EGF homology <EG2>

F;213-445/Domain: trypsin homology <TRY>

F;47-48-55-57-60-61-66-67-70-76/Modified site: gamma-carboxyglutamic acid (Glu) #status 1

F;112/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status predicted

F;121-130-139-150-146-159-161-174-182-320-239-255-373-387-398-426/Disulfide bonds: #stat

F;215-291-355/Binding site: carbohydrate (Asn) (covalent) #status predicted

F;254-300-402/Active site: His, Asp, Ser #status predicted

Query Match 66.7%; Score 132; DB 1; Length 461;
Best Local Similarity 56.8%; Pred. No. 2.9e-13;
Matches 25; Conservative 7; Mismatches 12; Indels 0; Gaps 0;

QY 1 ANSFLLXLRHSLRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44
DB 42 ANSFLEELRHSLEECIEECIECFEAEQIFQNVDDTLAFWSKH 85

RESULT 4

KXBO

Protein C (activated) (EC 3.4.21.69) precursor - bovine (fragment)

N;Alternate names: autoprothrombin IIA; plasma protein C

C;Species: Bos primigenius taurus (cattle)

C;Date: 30-Nov-1980 #sequence_revision 17-Mar-1987 #text_change 16-Jul-1999

C;Accession: A26250; A18385; A18386; A00928

R;Long, G.L.; Balagaje, R.M.; MacGillivray, R.T.A.

Proc. Natl. Acad. Sci. U.S.A. 81, 5653-5656, 1984

A;Title: Cloning and sequence of liver cDNA coding for bovine protein C.

A;Reference number: A26250; MUID:85014826; PMID:6091100

A;Accession: A26250

A;Molecule type: mRNA

A;Residues: 1-456 <LON>

R;Fernlund, P.; Stenflo, J.

A:Accession: S49075
A:Molecule type: mRNA
A:Residues: 1-482 <STA1>
A:Cross-references: EMBL:X79807; NID:G506600; PIDN:CAA56202.1; PID:G506601
A>Note: submitted to the EMBL Data Library, June 1994
A>Note: neither the complete nucleic acid sequence nor the complete translation are shown.
R:Stanton, C.; Rose, R.P.; Hutson, S.; Wallin, R.
Gene 169, 269-273, 1996

A:A>Title: Processing and expression of rat and human clotting factor-X encoding cDNAs.
A:Reference number: JG4670; MUID:96194815; PMID:8647460
A:Accession: JG4670
A:Molecule type: mRNA
A:Residues: 1-482 <STA2>
A:Cross-references: EMBL:X79807; NID:G506600; PIDN:CAA56202.1; PID:G506601
A:Experimental source: Cos-1 cell
R:Enyoji, K.; Miyazaki, K.; Kato, H.
J. Biochem. 109, 890-898, 1991

A:A>Title: Characterization of rat factors X and Xa: demonstration of factor Xa in rat plasma by immunoblotting.

A:Reference number: PS0190; MUID:92041742; PMID:1718949
A:Accession: PS0191
A:Molecule type: protein
A:Residues: 41-58, 'X', 60-65 <ENL1>
A:Accession: PS0190
A:Molecule type: protein
A:Residues: 183-186, 'X', 188-207 <ENJ2>
R:Murakawa, M.; Okamura, T.; Kamura, T.; Harada, M.; Niho, Y.
Eur. J. Haematol. 52, 162-168, 1994

A:A>Title: Analysis of the partial nucleotide sequences and deduced primary structures of tRNA^{Phe} from the Japanese quail (*Coturnix coturnix*).
A:Reference number: I46196; MUID:94221260; PMID:8168596
A:Accession: I62745
A>Status: preliminary; translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 295-383, 'G', 385-455 <MUR>
A:Cross-references: GB:D21215; NID:G415309; PIDN:BAA04756.1; PID:G455396
C:Function:
C:Description: catalyzes the proteolytic activation of prothrombin to thrombin in the presence of calcium ions.

A:Pathway: blood coagulation
C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglutamic acid
F:1-23/Domain: signal sequence #status predicted <SIG>
F:24-40/Domain: propeptide #status predicted <PRO>
F:25-84/Domain: Gla domain homology <GLA>
F:41-179/Product: coagulation factor X light chain #status predicted <LCH>
F:90-121/Domain: EGF homology <EG1>
F:129-164/Domain: EGF homology <EG2>
F:183-482/Product: coagulation factor X heavy chain #status predicted <HCH>
F:183-231/Domain: activation peptide #status predicted <APT>
F:232-482/Product: coagulation factor Xa heavy chain #status predicted <ACT>
F:232-480/Domain: trypsin homology <TRY>
F:46, 47, 54, 56, 59, 60, 65, 66, 69, 72, 79/Modified site: gamma-carboxyglutamic acid (Glu) #statu
F:57-62, 90-101, 95-110, 112-121, 129-140, 136-149, 151-164, 172-340, 238-243, 259-275, 388-402, 411
F:103/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status predicted
F:187/Binding site: carbonyl group (covalent) #status experimental
F:208/Binding site: carbonylate (Thr) (covalent) #status predicted
F:218/Binding site: carbonylate (Asn) (covalent) #status predicted
F:231-232/Cleavage site: Arg-Ile (coagulation factor IXa, coagulation factor VIIa) #statu
F:274, 320, 417/Active site: His, Asp, Ser #status predicted

Query Match 54.0%; Score 107; DB 1; Length 482;
Best Local Similarity 40.9%; Pred. No. 3.3e+09;
Matches 18; Conservative 9; Mismatches 17; Indels 0; Gaps 0;

OY 1 ANSFLXLKHSLSRPXCIXI:CDPFXAKXPEDVDDTLAPWSKH 44
||| : : : | : : : ||| : : : | : : :
DB 41 ANSFPEETIKGNLEECVEECSFEAREVFDEKTEFTFWNKY 84

RESULT 6
EXBO
coagulation factor Xa (EC 3.4.21.6) precursor - bovine
N:Alternate names: Stuart factor
C:Species: Bos primigenius taurus (cattle)
C>Date: 24-Apr-1984 #sequence revision 17-Mar-1987 #text change 16-Jul-1999

C:Accession: A22867; A14997; A12030; A34412; S39414; A00925
 Nucleic Acid Res. 12, 4481-4492, 1984
 A:Title: Blood coagulation factor X mRNA encodes a single polypeptide chain containing a
 A:Reference number: A22867; MUID:84247315; PMID:6330671
 A:Accession: A22867
 A:Molecule type: mRNA
 A:Residues: 1-487 <FUN>
 A:Cross-references: GB:X00673; NID:gl92; PIDN:CAA25286.1; PID:gl93
 R:Enfield, D.L.; Ericsson, L.H.; Fujikawa, K.; Walsh, K.A.; Neurath, H.; Titani, K.
 Biochemistry 19, 659-667, 1980
 A:Title: Amino acid sequence of the light chain of bovine factor X-1 (Stuart factor).
 A:Reference number: A14997; MUID:80130563; PMID:6766735
 A:Accession: A14997
 A:Molecule type: protein
 A:Residues: 41-102, 'N', 104-180 <ENF>
 R:McMullen, B.A.; Fujikawa, K.; Kisiel, W.
 Biochem. Biophys. Res. Commun. 115, 8-14, 1983
 A:Title: The occurrence of beta-hydroxyaspartic acid in the vitamin K-dependent blood coagulation factors.
 A:Reference number: A20274; MUID:83308813; PMID:6698526
 A:Contents: annotation; revision to residue 103
 R:Titani, K.; Fujikawa, K.; Enfield, D.L.; Ericsson, L.H.; Walsh, K.A.; Neurath, H.
 Proc. Natl. Acad. Sci. U.S.A. 72, 3082-3086, 1975
 A:Title: Bovine factor X-1 (Stuart factor): amino-acid sequence of heavy chain.
 A:Reference number: A12030; MUID:76053069; PMID:1059093
 A:Accession: A12030
 A:Molecule type: protein
 A:Residues: 183-292, 294-295, 'GDS', 299-334, 336-348, 'AE', 351-354, 356-441, 'GKFG', 446-492 <T>
 A:Note: Carbohydrate binding sites and disulfide bonds were determined
 R:Persson, E.; Selander, M.; Linse, S.; Drakenberg, T.; Oehlin, A.K.; Stenflo, J.
 J. Biol. Chem. 264, 16897-16904, 1989
 A:Title: Calcium binding to the isolated beta-hydroxyaspartic acid-containing epidermal
 A:Reference number: A34412; MUID:89380326; PMID:2799221
 A:Accession: A34412
 A:Molecule type: protein
 A:Residues: 85-126 <PER>
 A:Note: beta-hydroxyaspartic acid site
 R:Inoue, K.; Morita, T.
 Eur. J. Biochem. 218, 153-163, 1993
 A:Title: Identification of O-linked oligosaccharide chains in the activation peptides of
 A:Reference number: S39414; MUID:94062825; PMID:8243461
 A:Accession: S39414
 A:Molecule type: protein
 A:Residues: 183-196; 199-209; 216-233 <INO>
 A:Note: Carbohydrate binding sites
 R:Titani, K.; Hermanson, M.A.; Fujikawa, K.; Ericsson, L.H.; Walsh, K.A.; Neurath, H.; D
 Biochemistry 11, 4899-4903, 1972
 A:Title: Bovine factor X-1a (activated Stuart factor). Evidence of homology with mammalian
 A:Reference number: A12453; MUID:73053314; PMID:4264286
 A:Contents: annotation; active site
 R:Fujikawa, K.; Titani, K.; Davies, E.W.
 Proc. Natl. Acad. Sci. U.S.A. 72, 3359-3363, 1975
 A:Title: Activation of bovine factor X (Stuart factor): conversion of factor Xa/alpha to
 A:Reference number: A13504; MUID:76053121; PMID:1059122
 A:Contents: annotation; activation
 R:Suigo, T.; Bjork, I.; Holmgren, A.; Stenflo, J.
 J. Biol. Chem. 259, 5705-5710, 1984
 A:Title: Calcium-binding properties of bovine factor X lacking the gamma-carboxyglutamic
 A:Reference number: A38024; MUID:84185716; PMID:6546930
 A:Contents: annotation; calcium binding
 R:Morita, T.; Jackson, C.M.
 J. Biol. Chem. 261, 4008-4014, 1986
 A:Reference number: A38025; MUID:86140210; PMID:3949800
 A:Contents: annotation; sulfate binding
 A:Comment: Factor Xa converts prothrombin to thrombin during blood clotting.
 C:Comment: The two chains are formed from a single-chain precursor by the excision of two
 C:Comment: The activation peptide is cleaved by factor IXa (in the intrinsic pathway), C
 activation.
 C:Comment: Calcium binds to the gamma-carboxyglutamic acid (Gla) residues and, with str
 C:Comment: The gamma-carboxyglutamic acid residues arise by a posttranslational, vitamin
 C:Genetics:
 A:Gene: F10
 A:Map position: 13q34

C:Function:
 A:Description: catalyzes the proteolytic activation of prothrombin to thrombin in the pr
 A:Pathway: blood coagulation
 C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
 C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglutam
 F:1-15/Domain: signal sequence #status predicted <SIG>
 F:16-40/Domain: propeptide #status predicted <PRO>
 F:25-84/Domain: Gla domain homology <GLA>
 F:41-180/Product: coagulation factor X light chain #status experimental <LCH>
 F:90-121/Domain: EGF homology <EGF>
 F:129-164/Domain: EGF homology <EG2>
 F:183-492/Product: coagulation factor X heavy chain #status experimental <HCH>
 F:183-233/Domain: activation peptide #status experimental <APT>
 F:234-492/Product: coagulation factor Xa heavy chain #status experimental <AHC>
 F:234-461/Domain: trypsin homology <TRY>
 F:46 47 54 56 59 60 65 66 69 72 75 79/Modified site: gamma-carboxyglutamic acid (Glu) #s
 F:57-62, 90-101, 95-110, 112-121, 129-140, 136-149, 151-164, 172-341/Disulfide bonds: #status p
 F:103/Modified site: sulfuro-beta-hydroxyaspartic acid (Asp) #status experimental
 F:200/Binding site: tyrosine (Tyr) (covalent) (partial) #status experimental
 F:208, 485/Binding site: carboxylate (Thr) (covalent) #status experimental
 F:218/Binding site: carboxylate (Asn) (covalent) #status experimental
 F:233-234/Cleavage site: Arg-11e (coagulation factor IXa, coagulation factor VIIa) #stat
 F:240-245, 260-276, 389-403, 414-442/Disulfide bonds: #status experimental
 F:275, 321, 418/Active site: His, Asp, Ser #status predicted
 Query Match 52.0%; Score 103; DB 1; Length 492;
 Best Local Similarity 40.9%; Pred. No. 1.5e-08;
 Matches 18; Conservative 8; Mismatches 18; Indels 0; Gaps 0;
 QY 1 ANSLFXLRHSLSLRXCIXICDPFXKXKXIFEDVDVDTLAFWSKH 44
 Db 41 ANSLFEEVKQNLRECLERACSLSEAEAREVFEDAEQTFDEWSKY 84
 RESULT 7
 EXHU
 coagulation factor Xa (EC 3.4.21.6) precursor [validated] - human
 N:Alternate names: Stuart factor
 C:Species: Homo sapiens (man)
 C:Date: 15-Nov-1994 #sequence revision 02-May-1994 #text change 08-Dec-2000
 C:Accession: A24478; JQ0917; A2485; A25853; A22208; A21284; A20362; S39415; IS4051; A00'
 R:Leytus, S.P.; Foster, D.C.; Kurachi, K.; Davies, E.W.
 Biochemistry 25, 5098-5102, 1986
 A:Title: Gene for human factor X: a blood coagulation factor whose gene organization is
 A:Reference number: A24478; MUID:87026600; PMID:3768336
 A:Accession: A24478
 A:Molecule type: DNA
 A:Residues: 1-488 <LEV>
 A:Cross-references: GB:L29433; GB:M14327; NID:9459809; PIDN:AAA52764.1; PID:gl82831
 R:Messier, T.L.; Pittman, D.D.; Long, G.L.; Kaufman, R.J.; Church, W.R.
 Gene 99, 291-294, 1991
 A:Title: Cloning and expression in COS-1 cells of a full-length cDNA encoding human coag
 A:Reference number: JQ0917; MUID:91216473; PMID:1902434
 A:Accession: JQ0917
 A:Molecule type: mRNA
 A:Residues: 1-488 <MES>
 A:Cross-references: GB:M57285; NID:gl82389; PIDN:AAA52421.1; PID:gl82390
 R:Miao, C.H.; Leytus, S.P.; Chung, D.W.; Davie, E.W.
 J. Biol. Chem. 267, 7395-7401, 1992
 A:Title: Liver-specific expression of the gene coding for human factor X, a blood coagul
 A:Reference number: A2485; MUID:92218390; PMID:1313796
 A:Accession: A2485
 A:Molecule type: DNA
 A:Residues: 1-15 <MIA>
 A:Experimental source: liver
 A:Note: Sequence extracted from NCBI backbone (NCBI:93780, NCBI:93787)
 R:Kaul, R.K.; Hildebrand, B.; Roberts, S.; Jagadeeswaran, P.
 Gene 41, 311-314, 1986
 A:Title: Isolation and characterization of human blood-coagulation factor X cDNA.
 A:Reference number: A25853; MUID:86221713; PMID:3011603
 A:Accession: A25853
 A:Molecule type: mRNA
 A:Residues: 19-284, 'E', 289-488 <KAU>

A:Cross-references: GB:M22613; NID:g180335; PIDN:AAA51984.1; PID:g180336
F:Fung, M.R.; Hay, C.W.; Macgillivray, R.T.A.
Proc. Natl. Acad. Sci. U.S.A. 82, 3591-3595, 1985
A:Title: Characterization of an almost full-length cDNA coding for human blood coagulation factor VIIa
A:Reference number: A22208; MUID:85216545; PMID:2582420
A:Accession: A22208
A:Molecule type: mRNA
A:Residues: 13-441; S', 443-488 <F>
A:Cross-references: GB:X03194; NID:g182840; PIDN:AAA52490.1; PID:g182841
R:Levy, S.P.; Chung, D.W.; Kisiel, W.; Kurachi, K.; Davie, E.W.
Proc. Natl. Acad. Sci. U.S.A. 81, 3699-3702, 1984
A:Title: Characterization of a cDNA coding for human factor X
A:Reference number: A21284; MUID:84222036; PMID:6587384
A:Accession: A21284
A:Molecule type: mRNA
A:Residues: 13-284; B', 289-488 <LE2>
A:Cross-references: GB:X01886
R:McMullen, B.A.; Fujikawa, K.; Kisiel, W.; Sasagawa, T.; Howald, W.N.; Kwa, E.Y.; Weiss
Biochemistry 22, 2875-2884, 1983
A:Title: Complete amino acid sequence of the light chain of human blood coagulation factor VIIa
A:Reference number: A20362; MUID:83257207; PMID:6871167
A:Accession: A20362
A:Molecule type: protein
A:Residues: 41-179 <NCM>
R:Inoue, K.; Morita, T.
Eur. J. Biochem. 218, 153-163, 1993
A:Title: Identification of O-linked oligosaccharide chains in the activation peptides of human factor VIIa
A:Reference number: S39414; MUID:94062825; PMID:8243461
A:Accession: S39415
A:Molecule type: protein
A:Residues: 183-234 <INO>
A:Note: glycosylation sites
A:Note: identification and characterization of beta-hydroxyaspartic acid
R:Jagadeeswaran, P.; Reddy, S.V.; Rao, K.J.; Hamsabhusanam, K.; Lyman, G.
Gene 84, 517-519, 1989
A:Title: Cloning and characterization of the 5' end (exon 1) of the gene encoding human factor VIIa
A:Reference number: 154051; MUID:90128299; PMID:2612918
A:Accession: 154051
A:Status: translation not shown; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-23 <RES>
A:Cross-references: GB:M33297; NID:g183860; PIDN:AAA52636.1; PID:g553330
R:Padmanabhan, K.; Padmanabhan, K.P.; Tulinsky, A.; Park, C.H.; Bode, W.; Huber, R.; Bla
J. Mol. Biol. 232, 947-966, 1993
A:Title: Structure of human factor VIIa at 2.2 angstroms resolution.
A:Reference number: A49458; MUID:93360277; PMID:8355279
A:Contents: annotation; X-ray crystallography, 2.2 angstroms
C:Comment: The two chains held together by one disulfide bond are formed from a single-c
C:Function: The activation peptide is cleaved by factor IXa (in the intrinsic pathway) of
A:Gene: GDB:F10
A:Cross-references: GDB:119890; OMIM:227600
A:Map position: 13q34-13q34
A:Introns: 24/1; 77/3; 86/1; 124/1; 150/3; 249/3; 289/1
A:Note: deficiency of this factor causes Stuart disease
C:Function:
A:Description: catalyzes the proteolytic activation of prothrombin to thrombin in the pr
A:Pathway: blood coagulation
C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxylglut
F:1-23/Domain: signal sequence #status predicted <SIG>
F:24-40/Domain: propeptide #status predicted <PRO>
F:25-84/Domain: Gla domain homology <GLA>
F:41-179/Product: coagulation factor X light chain #status experimental <LCH>
F:90-121/Domain: EGF homology <EGF>
F:129-164/Domain: EGF homology <EG2>
F:183-498/Product: coagulation factor X heavy chain #status experimental <HCH>
F:183-234/Domain: activation peptide #status experimental <APT>
F:235-498/Product: coagulation factor Xa heavy chain #status experimental <ACT>
F:235-462/Domain: trypsin homology <TRY>
F:46-47/Domain: gamma-carboxylglutamic acid (Glu) #stat
F:57-62/Disulfide bonds: #status predicted
F:90-101,95-110,112-121,129-140,136-149,151-164,172-342,241-246,261-277,390-404,415-443/

F:103/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status experimental
F:139,211/Binding site: carbohydrate (Thr) (covalent) #status experimental
F:221,231/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:234-235/Cleavage site: Arg-Tle (coagulation factor IXa, coagulation factor VIIa) #statu
F:276,322,419/Active site: His, Asp, Ser #status experimental
Query Match 51.5%; Score 102; DB 1; Length 488;
Best Local Similarity 40.9%; Pred. No. 2.1e-08;
Matches 18; Conservative 8; Mismatches 18; Indels 0; Gaps 0;
QY 1 ANSFLXXLRHSSLRXCIXXICDFFXXAXXIFEDVDDTLAFWSKH 44
DB 41 ANSFLXEMKKHLEKRCMRETCSEYERARVFDSDXTNEFWNKY 84

RESULT 8

I46932
C:coagulation factor VII - rabbit
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 04-Sep-1997 #sequence_revision 04-Sep-1997 #text_change 12-Feb-1999
A:Accession: I46932
R:Brothers, A.B.; Clarke, B.J.; Sheffield, W.P.; Blajchman, M.A.
Thromb. Res. 69, 231-238, 1993
A:Title: Complete nucleotide sequence of the cDNA encoding rabbit coagulation factor VII.
A:Reference number: I46932; MUID:93190306; PMID:8383365
A:Accession: I46932
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-443 <ERO>
A:Cross-references: GB:S56300; NID:g266294; PID:g266295
C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
F:24-83/Domain: Gla domain homology <GLA>
F:89-120/Domain: EGF homology <EG1>
F:130-166/Domain: EGF homology <EG2>
F:192-425/Domain: trypsin homology <TRY>

Query Match

47.5%; Score 94; DB 2; Length 443;
Best Local Similarity 43.9%; Pred. No. 3.7e-07;
Matches 18; Conservative 5; Mismatches 18; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHSSLRXCIXXICDFFXXAXXIFEDVDDTLAFW 41

DB 40 ANSFLXELRPGSLERECKELCSFEAREVFQSTERTKQFW 80

RESULT 9

KRFU7
C:coagulation factor VIIa (EC 3.4.21.21) precursor [validated] - human
C:Species: Homo sapiens (man)
C:Date: 19-May-1989 #sequence_revision 19-May-1994 #text_change 08-Dec-2000
A:Accession: A28322; A23819; A31186; B31186; S63524
R:O'Hara, P.J.; Grant, P.J.; Haldeman, B.A.; Gray, C.L.; Insley, M.Y.; Hagen, F.S.; Murza
Proc. Natl. Acad. Sci. U.S.A. 84, 5158-5162, 1987
A:Title: Nucleotide sequence of the gene coding for human factor VII.
A:Reference number: A28322; MUID:87260948; PMID:3037537
A:Accession: A28322
A:Molecule type: DNA
A:Residues: 1-466 <OHA>
A:Cross-references: GB:J02933; NID:g180333; PIDN:AAA51983.1; PID:g180334
R:Hagen, F.S.; Gray, C.L.; O'Hara, P.; Grant, P.J.; Saari, G.C.; Woodbury, R.G.; Hart, C
Proc. Natl. Acad. Sci. U.S.A. 83, 2412-2416, 1986
A:Title: Characterization of a cDNA coding for human factor VII.
A:Reference number: A23819; MUID:86205965; PMID:3486420
A:Accession: A23819
A:Molecule type: mRNA
A:Residues: 1-466 <HAG>
A:Cross-references: GB:M13232; NID:g182799; PIDN:AAA8040.1; PID:g182801
R:Thim, L.; Bicorn, S.; Christensen, M.; Nicolaisen, E.M.; Lund-Hansen, T.; Pedersen, A.;
Biochemistry 27, 7795-7793, 1988
A:Title: Amino acid sequence and posttranslational modifications of human factor VII-a f;
A:Reference number: A90539; MUID:89088153; PMID:3264725
A:Accession: A31186
A:Molecule type: protein

A;Residues: 61-212 <TH1>
A;Accession: B31186
A;Molecule type: protein
R;Biojorn, S.; Foster, D.C.; Thim, L.; Wiberg, F.C.; Christensen, M.; Komiyama, Y.; Pedersen, J. Biol. Chem. 266, 11051-11057, 1991
A;Title: Human plasma and recombinant factor VII. Characterization of O-glycosylations and A;Reference number: A40529; MUID:91250411; PMID:1904059
A;Contents: annotation; carbohydrate binding sites
R;Persson, E.; Petersen, L.C.
Eur. J. Biochem. 234, 293-300, 1995
A;Title: Structurally and functionally distinct Ca(2+) binding sites in the gamma-carboxy A;Reference number: S63524; MUID:96096752; PMID:8529655
A;Accession: S63524
A;Molecule type: protein
A;Residues: 61-65;99-103;105-109;213-217;308-312 <PER>
C;Genetics:
A;Gene: GDB:F7
A;Cross-references: GDB:1119897; OMIM:227500
A;Map position: 13q34-13q34
A;Introns: 22/1; 44/1; 97/3; 106/1; 144/1; 191/1; 227/3; 269/1
C;Function:
A;Description: catalyzes the proteolytic activation of coagulation factor X in the presence of coagulation factor IX in the presence of calcium and tissue factor
A;Pathway: blood coagulation extrinsic pathway
C;Superfamily: coagulation factor X; EGF homology
C;Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglutamic acid
F;1-20/Domain: signal sequence #status predicted <SIG>
F;21-60/Domain: propeptide #status predicted <PRO>
F;45-104/Domain: Gla domain homology <GLA>
F;61-141/Product: coagulation factor VIIa light chain #status experimental <VAL>
F;110-141/Domain: EGF homology <EG1>
F;151-187/Domain: EGF homology <EG2>
F;213-466/Product: coagulation factor VIIa heavy chain #status experimental <MA2>
F;213-447/Domain: trypsin homology <TRY>
F;66-67;74-76;80-85;86-89;95/Modified site: gamma-carboxyglutamic acid (Glu) #status predicted <SIG>
F;77-82;110-121;115-130;132-141;151-162;158-172;174-187;195-322;219-224;238-254;370-389;1120/Binding site: carboxylate (Ser) (covalent) #status experimental
F;123/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status absent
F;205;382/Binding site: Arg-Arg (covalent) (Asn) (covalent) #status experimental
F;212-213/Cleavage site: Arg-Ile (coagulation factor XIIIa) #status experimental
F;253;302;404/Active site: His, Asp, Ser #status predicted
F;350-351/Cleavage site: Arg-Gly (coagulation factor Xa) #status predicted
Query Match 46.5%; Score 92; DB 1; Length 466;
Best Local Similarity 46.3%; Pred. No. 8.3e-07;
Matches 19; Conservative 4; Mismatches 18; Indels 0; Gaps 0;
QY 1 ANSFLXLRHSSLRXCIXXICDFXXAKXIFEDVDDTLAFW 41
DB 61 ANAFLEELRPGSLRCKEVEQCSFEEREIFKDAERTKLEW 101
RESULT 10
thrombin (EC 3.4.21.5) precursor - rat
C;Species: Rattus norvegicus (Norway rat)
C;Date: 07-May-1993 #sequence revision 07-May-1993 #text_change 03-May-2002
C;Accession: S10511; A60576; B42696
R;Dihantich, M.; Monard, D.
Nucleic Acids Res. 18, 4251, 1990
A;Title: cDNA sequence of rat prothrombin.
A;Reference number: S10511; MUID:90332426; PMID:2377469
A;Accession: S10511
A;Molecule type: mRNA
A;Residues: 1-617 <DIH>
A;Cross-references: ENBL:X52835; NID:956969; PIDN:CAA37017.1; PID:956970
R;Henrikson, K.P.; Jasin, E.E.; Greenwood, J.A.; Dickerman, H.W.
Endocrinology 126, 167-175, 1990
A;Title: Prothrombin levels are increased in the estrogen-treated immature rat uterus.
A;Reference number: A60576; MUID:90091942; PMID:2293980
A;Accession: A60576
A;Molecule type: protein

A;Residues: 44-58 <HEN>
A;Note: the authors purified the proenzyme from the estrogen-stimulated maturing rat ute.
R;Banfield, D.K.; MacGillivray, R.T.A.
Proc. Natl. Acad. Sci. U.S.A. 89, 2779-2783, 1992
A;Title: Partial characterization of vertebrate prothrombin cDNAs: amplification and seq
A;Reference number: A42696; MUID:92212913; PMID:1557383
A;Accession: B42696
A;Status: Preliminary
A;Molecule type: mRNA
A;Residues: 383-617; E' <BAN>
A;Cross-references: GB:M81397
C;Superfamily: thrombin; Gla domain homology; kringle homology; trypsin homology
C;Keywords: blood coagulation; calcium binding; carboxyglutamic acid; glycoprotein; hydr
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-43/Domain: propeptide #status predicted <PRO>
F;28-88/Domain: Gla domain homology <GLA>
F;44-617/Product: prothrombin #status experimental <PMAT>
F;109-187/Domain: kringle homology <KRI>
F;215-292/Domain: kringle homology <KR2>
F;360-609/Domain: trypsin homology <TRY>
F;50-51;58;60;63;64;69;70;73;76/Modified site: gamma-carboxyglutamic acid (Glu) #status
F;61-66;91-104;109-187;130-170;158-182;215-292;236-276;264-287;332-478;387-403;532-546;5
F;402;458;564/Active site: His, Asp, Ser #status predicted
Query Match 39.6%; Score 78.5; DB 2; Length 617;
Best Local Similarity 40.0%; Pred. No. 0.00017;
Matches 18; Conservative 5; Mismatches 21; Indels 1; Gaps 1;
QY 1 ANSFLXLRHSSLRXCIXXICDFXXAKXIFEDVDDTLAFWSKH 44
DB 44 ANSGFLEELRKGNLRECVQCSFEERFALESPQDDTDFWAKY 88
RESULT 11
thrombin (EC 3.4.21.5) precursor - mouse
C;Species: Mus musculus (house mouse)
C;Date: 14-Dec-1990 #sequence revision 14-Dec-1990 #text_change 03-May-2002
C;Accession: A35827; A42696; S12081
R;Degen, S.J.F.; Schaefer, L.A.; Jamison, C.S.; Grant, S.G.; Fitzgibbon, J.J.; Pai, J.A.
DNA Cell Biol. 9, 487-498, 1990
A;Title: Characterization of the cDNA coding for mouse prothrombin and localization of t
A;Reference number: A35827; MUID:91025551; PMID:2222810
A;Accession: A35827
A;Status: Preliminary
A;Molecule type: mRNA
A;Residues: 1-618 <DEG>
A;Cross-references: GB:X52308; NID:953813; PIDN:CAA36548.1; PID:953814
A;Experimental source: strain C57BL/6
A;Note: the data were obtained from females resulting from the cross of M. domesticus an
R;Banfield, D.K.; MacGillivray, R.T.A.
Proc. Natl. Acad. Sci. U.S.A. 89, 2779-2783, 1992
A;Title: Partial characterization of vertebrate prothrombin cDNAs: amplification and seq
A;Reference number: A42696; MUID:92212913; PMID:1557383
A;Accession: A42696
A;Status: Preliminary
A;Molecule type: mRNA
A;Residues: 384-618; E' <BAN>
A;Cross-references: GB:M81394
C;Superfamily: thrombin; Gla domain homology; kringle homology; trypsin homology
C;Keywords: blood coagulation; calcium binding; carboxyglutamic acid; glycoprotein; hydr
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-43/Domain: propeptide #status predicted <PRO>
F;28-88/Domain: Gla domain homology <GLA>
F;44-618/Product: prothrombin B #status predicted <MAT>
F;109-187/Domain: kringle homology <KRI>
F;215-293/Domain: kringle homology <KR2>
F;361-610/Domain: trypsin homology <TRY>
F;50-51;58;60;63;64;69;70;73;76/Modified site: gamma-carboxyglutamic acid (Glu) #status
F;61-66;91-104;109-187;130-170;158-182;215-293;236-276;264-288;333-479;388-404;533-547;5
F;403;459;565/Active site: His, Asp, Ser #status predicted
Query Match 39.6%; Score 78.5; DB 2; Length 618;

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OM protein - protein search, using sw model

Run on: March 1, 2004, 09:58:27 ; Search time 10.5 Seconds
(without alignments)
218.199 Million cell updates/sec

Title: SEQ1-32GLU-33ASP

Perfect score: 198

Sequence: 1 ANSFLXLRHSSLRXCIXX.....XXAKXITVDVDTLAFWSKH 44

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	172	86.9	461	1	PRTC_HUMAN
2	141	71.2	458	1	PRTC_RABIT
3	133	67.2	461	1	PRTC_MOUSE
4	132	66.7	461	1	PRTC_RAT
5	126	63.6	459	1	PRTC_PIG
6	115	58.1	456	1	PRTC_BOVIN
7	103	52.0	492	1	FA10_BOVIN
8	102	51.5	488	1	FA10_HUMAN
9	96	48.5	231	1	TWG3_HUMAN
10	95	48.0	490	1	FA10_RABIT
11	94	47.5	444	1	FA10_HOPST
12	92	46.5	466	1	FA10_HUMAN
13	81	40.9	218	1	TWG1_HUMAN
14	78.5	39.6	617	1	THRB_RAT
15	78.5	39.6	618	1	THRB_MOUSE
16	78	39.4	226	1	TWG4_HUMAN
17	78	39.4	407	1	FA10_BOVIN
18	77	38.9	376	1	FA10_HOPST
19	76	38.4	622	1	THRB_HUMAN
20	75	37.9	376	1	FA10_TROCA
21	75	37.9	475	1	FA10_CHICK
22	74	37.4	649	1	PRPS_VACUO
23	74	37.4	676	1	PRPS_HUMAN
24	73	36.9	446	1	FA10_MOUSE
25	73	36.9	452	1	FA10_CANFA
26	73	36.9	459	1	FA10_MOUSE
27	73	36.9	646	1	PRPS_RABIT
28	72	36.4	675	1	PRPS_BOVIN
29	70	35.4	675	1	PRPS_RAT
30	69	34.8	461	1	FA10_HUMAN
31	69	34.8	461	1	FA10_PANTR
32	66	33.3	416	1	FA10_BOVIN
33	65	32.8	98	1	FA10_NOTCS

ALIGNMENTS

RESULT 1

ID	PRTC_HUMAN	STANDARD	PRT	461 AA
AC	P04070: Q15189; Q15190; Q16001;			
DT	01-NOV-1986 (Rel. 03, Created)			
DT	01-NOV-1986 (Rel. 03, Last sequence update)			
DT	15-MAR-2004 (Rel. 43, Last annotation update)			
DE	Vitamin-K-dependent protein C precursor (EC 3.4.21.69)			
DE	(Autoproteolysis IIA) (Anticoagulant protein C) (Blood coagulation factor XIV)			
GN	PROC.			
OS	Homo sapiens (Human)			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Theria; Primates; Catarrhini; Hominoidea; Homo.			
OX	NCBI_TaxID=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=85270390; PubMed=2991887;			
RA	Forster D.C., Yonitake S., Davie E.W.;			
RT	"The nucleotide sequence of the gene for human protein C.";			
RL	Proc. Natl. Acad. Sci. U.S.A. 82:4673-4677(1985).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=85269639; PubMed=2991859;			
RA	Beckmann R.J., Schmidt R.J., Santerre R.F., Plutsky J., Crabtree G.R.,			
RA	Long G.L.;			
RT	"The structure and evolution of a 461 amino acid human protein C			
RT	precursor and its messenger RNA, based upon the DNA sequence of			
RT	cloned human liver cDNAs.";			
RL	Nucleic Acids Res. 13:5233-5247(1985).			
RN	[3]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=86120978; PubMed=3511471;			
RA	Plutsky J., Hoskins J.A., Long G.L., Crabtree G.R.;			
RT	"Evolution and organization of the human protein C gene.";			
RL	Proc. Natl. Acad. Sci. U.S.A. 83:546-550(1986).			
RN	[4]			
RP	SEQUENCE FROM N.A.			
RA	Rieder M.J., Carrington D.P., Chung M.-W., Lee K.L., Poel C.L., Yi Q.,			
RA	Nickerson D.A.;			
RL	Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.			
RN	[5]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Colon;			
RX	MEDLINE=22338237; PubMed=12477932;			
RA	Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,			
RA	Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,			
RA	Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,			
RA	Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,			
RA	Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,			
RA	Stapleton L., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,			
RA	Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C.,			
RA	Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,			
RA	Bosak S.A., McSwan P.O., McKernan K.J., Malek J.A., Gunaratne P.H.,			
RA	Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,			
RA	Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,			

34	64	32.3	625	1	THRB_BOVIN
35	63	31.8	675	1	PRTC_MOUSE
36	61.5	31.1	396	1	PRTC_BOVIN
37	58	29.3	202	1	TWG2_HUMAN
38	57.5	29.0	400	1	PRTC_HUMAN
39	53	26.8	730	1	PRIA_HAIEIN
40	51	25.8	1363	1	VGR3_MOUSE
41	50	25.3	501	1	MKCI_CANAL
42	50	25.3	1298	1	VGR3_HUMAN
43	49.5	25.0	941	1	VP11_HUMAN
44	49.5	25.0	941	1	VP11_MOUSE
45	49	24.7	1343	1	VGR2_RAT

P00735	bos taurus
Q08761	mus musculus
P00744	bos taurus
O14669	homo sapien
P22891	homo sapien
P44647	haemophilus
P35917	mus musculus
P43058	candida alb
P35916	homo sapien
Q9h270	homo sapien
Q91W86	mus musculus
O08775	rattus norv

RA Fahey J., Helton E., Kettaman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Granwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smallus D.E.,
RA Schnerch A., Schein J.B., Jones S.J.M., Marra M.A.,
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [16]
RP SEQUENCE OF 106-461 FROM N.A.
RX MEDLINE=84272714; PubMed=6589623;
RA Foster D.C., Davie E.W.;
RT "Characterization of a cDNA coding for human protein C.";
RN Proc. Natl. Acad. Sci. U.S.A. 81:4766-4770(1984).
RN [7]
RP CARBOHYDRATE-LINKAGE SITE ASN-371.
RX MEDLINE=90293094; PubMed=1694179;
RA Mileteich J.P., Broze G.J. Jr.;
RT "Beta protein C is not glycosylated at asparagine 329. The rate of
RT translation may influence the frequency of usage at asparagine-X-
RT cysteine sites.";
RL J. Biol. Chem. 265:11397-11404(1990).
RN [8]
RP HYDROXYLATION.
RX MEDLINE=92184750; PubMed=1544894;
RA Harris R.J., Ling V.T., Spellman M.W.;
RT "O-linked fucose is present in the first epidermal growth factor
RT domain of factor XII but not protein C.";
RL J. Biol. Chem. 267:5102-5107(1992).
RN [9]
RP 3D-STRUCTURE MODELING OF 175-450.
RX MEDLINE=94272342; PubMed=8003977;
RA Fisher C.I., Greengard J.S., Griffin J.H.;
RT "Models of the serine protease domain of the human antithrombotic
RT plasma factor activated protein C and its zymogen.";
RL Protein Sci. 3:588-599(1994).
RN [10]
RP X-RAY CRYSTALLOGRAPHY (2.8 ANGSTROMS) OF 84-461.
RX MEDLINE=97157472; PubMed=9003757;
RA Mather T., Oganessyan V., Hof P., Huber R., Foundling S., Esmon C.,
RA Bode W.;
RT "The 2.8 A crystal structure of Gla-domainless activated protein C.";
RL EMBO J. 15:6822-6831(1996).
RN [11]
RP REVIEW ON PROC VARIANTS.
RX MEDLINE=93190290; PubMed=8446940;
RA Reitsma P.H., Poort S.R., Bernardi F., Gandrille S., Long G.L.,
RA Sala N., Cooper D.N.;
RT "Protein C deficiency: a database of mutations. For the Protein C & S
RT Subcommittee of the Scientific and Standardization Committee of the
RT International Society on Thrombosis and Haemostasis.";
RN Thromb. Haemost. 69:177-84(1993).
RN [12]
RP VARIANT PROC DEFICIENCY CYS-444.
RX MEDLINE=87204221; PubMed=2437584;
RA Romeo G., Hassan H.J., Staemfli S., Roncuzzi L., Cianetti L.,
RA Leonardi A., Vicente V., Mannucci P.M., Bertina R.M., Peschle C.,
RA Cortese R.;
RT "Hereditary thrombophilia: identification of nonsense and missense
RT mutations in the protein C gene.";
RL Proc. Natl. Acad. Sci. U.S.A. 84:2829-2832(1987).
RN [13]
RP VARIANT PROC DEFICIENCY TRP-211.
RX MEDLINE=90098906; PubMed=2602169;
RA Grundy C.B., Chitcolle A., Talbot S., Bevan D., Kakkar V.V.,
RA Cooper D.N.;
RT "Protein C London 1: recurrent mutation at Arg-169 (GGG-->TGG) in
RT the protein C gene causing thrombosis.";
RL Nucleic Acids Res. 17:10513-10513(1989).
RN [14]
RP VARIANT PROC DEFICIENCY CYS-272.
RX MEDLINE=91329836; PubMed=1868249;
RA Reitsma P.H., Poort S.R., Allaart C.F., Briet E., Bertina R.M.;
RT "The spectrum of genetic defects in a panel of 40 Dutch families with
RT symptomatic protein C deficiency type I: heterogeneity and founder
RT effects.";
RL Blood 78:890-894(1991).
RN [15]
RP VARIANTS PROC DEFICIENCY ALA-62 AND MET-76.
RX MEDLINE=92190481; PubMed=1347706;
RA Bovill E.G., Tomczak J.A., Grant B., Bhushan F., Pillemer E.,
RA Rainville I.R., Long G.L.;
RT "Protein C variant: symptomatic type II protein C deficiency
RT associated with two GLA domain mutations.";
RL Blood 79:1456-1465(1992).
RN [16]
RP VARIANT PROC DEFICIENCY ASP-418.
RX MEDLINE=92305321; PubMed=1611081;
RA Sugahara Y., Miura O., Yuen P., Aoki N.;
RT "Protein C deficiency Hong Kong 1 and 2: hereditary protein C
RT deficiency caused by two mutant alleles, a 5-nucleotide deletion and
RT a missense mutation.";
RL Blood 80:126-133(1992).
RN [17]
RP VARIANT PROC DEFICIENCY LEU-289.
RX MEDLINE=92380860; PubMed=1511988;
RA Grundy C.B., Chisholm M., Kakkar V.V., Cooper D.N.;
RT "A novel homozygous missense mutation in the protein C (PROC) gene
RT causing recurrent venous thrombosis.";
RL Hum. Genet. 89:683-684(1992).
RN [18]
RP VARIANTS PROC DEFICIENCY GLN-220 AND TRP-220.
RX MEDLINE=92380861; PubMed=1511989;
RA Grundy C.B., Schulman S., Tengborn L., Kakkar V.V., Cooper D.N.;
RT "Two different missense mutations at Arg 178 of the protein C (PROC)
RT gene causing recurrent venous thrombosis.";
RL Hum. Genet. 89:685-686(1992).
RN [19]
RP VARIANT PROC DEFICIENCY GLN-220.
RX MEDLINE=93250852; PubMed=1301959;
RA Gandrille S., Vidaud M., Alach M., Alhenc-Gelas M., Fischer A.M.,
RA Couault-Heilmann M., Toulon P., Flessinger J.N., Goossens M.;
RT "Two novel mutations responsible for hereditary type I protein C
RT deficiency: characterization by denaturing gradient gel
RT electrophoresis.";
RL Hum. Mutat. 1:491-500(1992).
RN [20]
RP VARIANT PROC DEFICIENCY SER-334.
RX MEDLINE=92276939; PubMed=1593215;
RA Yamamoto K., Matsushita T., Sugiura I., Takamatsu J., Iwasaki E.,
RA Wada H., Deguchi K., Shirakawa S., Saito H.;
RT "Homozygous protein C deficiency: identification of a novel missense
RT mutation that causes impaired secretion of the mutant protein C.";
RL J. Lab. Clin. Med. 119:682-689(1992).
RN [21]
RP VARIANTS PROC DEFICIENCY TRP-38; CYS-42; HIS-42; GLN-271 AND ASN-294.
RX MEDLINE=93313192; PubMed=8324221;
RA Gandrille S., Alhenc-Gelas M., Gaussem P., Aillaud M.-F., Dupuy E.,
RA Juhan-Vague I., Alach M.;
RT "Five novel mutations located in exons III and IX of the protein C
RT gene in patients presenting with defective protein C anticoagulant
RT activity.";
RL Blood 82:159-168(1993).
RN [22]
RP VARIANTS PROC DEFICIENCY GLY-14; GLN-211; TVR-244; GLN-253; LEU-321;
RX CYS-328; ILE-385; THR-388 AND VAL-388.
RA MEDLINE=93271391; PubMed=8499565;
RA Poort S.R., Pabinger-Fasching I., Mannhalter C., Reitsma P.H.,
RA Bertina R.M.;
RT "Twelve novel and two recurrent mutations in 14 Austrian families
RT with hereditary protein C deficiency.";
RL Blood Coagul. Fibrinolysis 4:273-280(1993).
RN [23]
RP VARIANT PROC DEFICIENCY TRP-57.
RX MEDLINE=93271396; PubMed=8499568;

Query Match 86.9%; Score 172; DB 1; Length 461;
Best Local Similarity 75.0%; Pred. No. 5.9e-21;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXHLHSHSLXRCXIXICDFFXAXKXIFEDVDTLAFWSKH 44
DB 43 ANSFLXHLHSHSLXRCXIXICDFFXAXKXIFEDVDTLAFWSKH 86

RESULT 2

PTIC RABIT
ID PRTC RABIT STANDARD; PRT; 458 AA.
AC Q28661;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Vitamin-K-dependent protein C precursor (EC 3.4.21.69)
DE (Autoproteolytic cleavage of protein C)
DE factor XIV (Fragment).
GN PROC.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]
RF SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Shen L., He X., Dahlback B.;
RL Submitted (FEB-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Protein C is a vitamin K-dependent serine protease that
CC regulates blood coagulation by inactivating factors Va and VIIIa
CC in the presence of calcium ions and phospholipids.
CC -!- CATALYTIC ACTIVITY: Degradation of blood coagulation factors Va
CC and VIIIa.
CC -!- SUBUNIT: Synthesized as a single chain precursor, which is cleaved
CC into a light chain and a heavy chain held together by a disulfide
CC bond. The enzyme is then activated by thrombin, which cleaves a
CC tetradecapeptide from the amino end of the heavy chain; this
CC reaction, which occurs at the surface of endothelial cells, is
CC strongly promoted by thrombomodulin.
CC -!- TISSUE SPECIFICITY: Plasma; synthesized in the liver.
CC -!- PTM: The vitamin K-dependent, enzymatic carboxylation of some Glu
CC residues allows the modified protein to bind calcium.
CC -!- MISCELLANEOUS: Calcium also binds, with stronger affinity to
CC another site, beyond the GLA domain. This GLA-independent binding
CC site is necessary for the recognition of the thrombin-
CC thrombomodulin complex.
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 EGF-like domains.
CC
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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (see <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
CC
CC EMBL; U49933; AAA92956.1;
CC HSSP; P04070; 1PCU.
CC
CC MEROPS; S01.218; -;
CC InterPro; IPR000152; Asx_hydroxyl_S.
CC InterPro; IPR009003; Cys_Ser_trypsin.
CC InterPro; IPR001881; EGF_Ca.
CC InterPro; IPR006209; EGF_Like.
CC InterPro; IPR002383; GLA blood.
CC InterPro; IPR006210; IEGF.
CC InterPro; IPR001254; Peptidase_S1.
CC InterPro; IPR001314; Peptidase_S1A.
CC InterPro; IPR000294; VitK_dep_GLA.
CC Pfam; PF00008; EGF; 2.
CC Pfam; PF00594; Gla; 1.

Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00181; EGF; 2.
DR SMART; SM00069; GLA; 1.
DR SMART; SM00020; TYD_SPC; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS00026; EGF_3; 1.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
KW Blood coagulation; Glycoprotein; Serine protease;
KW Gamma-carboxyglutamic acid; Calcium-binding; Vitamin K; Hydroxylation;
KW EGF-like domain; Repeat; Endothelial cell; Hydrolase; Signal.
FT NON_TER 1 1
FT SIGNAL <1 27 BY SIMILARITY.
FT PROPEP 28 36 BY SIMILARITY.
FT CHAIN 37 458 VITAMIN K-DEPENDENT PROTEIN C.
FT CHAIN 37 192 PROTEIN C LIGHT CHAIN (BY SIMILARITY).
FT CHAIN 195 458 PROTEIN C HEAVY CHAIN (BY SIMILARITY).
FT PEPTIDE 195 209 ACTIVATION PEPTIDE (BY SIMILARITY).
FT SITE 209 210 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
FT DOMAIN 91 126 EGF-LIKE 1.
FT DOMAIN 130 170 EGF-LIKE 2.
FT DOMAIN 210 458 SERINE PROTEASE.
FT MOD_RES 42 42 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 43 43 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 50 50 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 52 52 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 55 55 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 56 56 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 61 61 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 62 62 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 65 65 GAMMA-CARBOXYGLUTAMIC ACID
(BY SIMILARITY).
FT MOD_RES 107 107 HYDROXYLATION (BY SIMILARITY).
FT ACT_SITE 250 250 CHARGE RELAY SYSTEM.
FT ACT_SITE 296 296 CHARGE RELAY SYSTEM.
FT ACT_SITE 399 399 CHARGE RELAY SYSTEM.
FT DISULFID 53 58 BY SIMILARITY.
FT DISULFID 86 105 BY SIMILARITY.
FT DISULFID 95 100 BY SIMILARITY.
FT DISULFID 99 114 BY SIMILARITY.
FT DISULFID 116 125 BY SIMILARITY.
FT DISULFID 134 145 BY SIMILARITY.
FT DISULFID 141 154 BY SIMILARITY.
FT DISULFID 156 169 BY SIMILARITY.
FT DISULFID 177 316 INTERCHAIN (BY SIMILARITY).
FT DISULFID 235 251 BY SIMILARITY.
FT DISULFID 370 384 BY SIMILARITY.
FT DISULFID 395 423 BY SIMILARITY.
FT CARBOHYD 133 133 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 287 287 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 352 352 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 458 AA; 51087 MW; D75A5F990C8F29D7 CRC64;

Query Match 71.2%; Score 141; DB 1; Length 458;
Best Local Similarity 61.4%; Pred. No. 9e-16;
Matches 27; Conservative 4; Mismatches 13; Indels 0; Gaps 0;

QY 1 ANSFLXHLHSHSLXRCXIXICDFFXAXKXIFEDVDTLAFWSKH 44


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FT DISULFID 396 424 BY SIMILARITY
FT CARBOHYD 138 138 N-LINKED (GLCNAC. . .) (POTENTIAL)
FT CARBOHYD 292 292 N-LINKED (GLCNAC. . .) (POTENTIAL)
FT CARBOHYD 353 353 N-LINKED (GLCNAC. . .) (POTENTIAL)
SQ SEQUENCE 459 AA; 51866 MW; 8541AAC14CC16D09 CRC64;

Query Match 63.6%; Score 126; DB 1; Length 459;
Best Local Similarity 54.5%; Pred. No. 2,9e-13;
Matches 24; Conservative 7; Mismatches 13; Indels 0; Gaps 0;

QY 1 ANSFLXKHLHSSLRXCIXXICDFAKXKXIFEDVDVDTAFWSKH 44
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 42 ANSFLFELRPPSLERCKEETCDFEARBIQONTENTAFWSKY 85

RESULT 6
PRTC_BOVIN STANDARD; PRT; 456 AA.
AC P00745;
DT 21-JUL-1986 (Rel. 01, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Vitamin-K-dependent protein C precursor (EC 3.4.21.69)
DE (Autoprothrombin IIA) (Anticoagulant protein C) (Blood coagulation
DE factor XIV) (Fragment).
GN PROC.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCB_TaxID:9913;
[1]
SEQUENCE FROM N.A.
RX MEDLINE=85014826; PubMed=6091100;
RA Long G.L., Balagaje R.M., McGillivray R.T.A.;
RT "Cloning and sequencing of liver cDNA coding for bovine protein C.";
RL "Proc. Natl. Acad. Sci. U.S.A. 81:5653-5656(1984)".
[2]
SEQUENCE OF 40-194, AND CARBOHYDRATE-LINKAGE SITE ASN-136.
RX MEDLINE=83007325; PubMed=6896876;
RA Fernlund P., Stenflo J.;
RT "Amino acid sequence of the light chain of bovine protein C.";
RL J. Biol. Chem. 257:12170-12179(1982).
[3]
REVISION TO 110.
RX MEDLINE=83169769; PubMed=6572939;
RA Drakenberg T., Fernlund P., Roepstorff P., Stenflo J.;
RT "Beta-hydroxyaspartic acid in vitamin K-dependent protein C.";
RL "Proc. Natl. Acad. Sci. U.S.A. 80:1802-1806(1983)".
[4]
SEQUENCE OF 197-456, AND CARBOHYDRATE-LINKAGE SITES ASN-289; ASN-350
AND ASN-366.
RX MEDLINE=83007326; PubMed=6896877;
RA Stenflo J., Fernlund P.;
RT "Amino acid sequence of the heavy chain of bovine protein C.";
RL J. Biol. Chem. 257:12180-12190(1982).
[5]
PROCESSING, AND CALCIUM-BINDING DATA.
RX MEDLINE=83213513; PubMed=6304092;
RA Esmon N.L., Debault L.E., Esmon C.T.;
RT "Proteolytic formation and properties of gamma-carboxyglutamic acid-
domainless protein C.";
RL J. Biol. Chem. 258:5548-5553(1983).
[6]
PROCESSING, AND CALCIUM-BINDING DATA.
RX MEDLINE=83213514; PubMed=6406503;
RA Johnson A.B., Esmon N.L., Laue T.M., Esmon C.T.;
RT "Structural changes required for activation of protein C are induced
by Ca2+ binding to a high affinity site that does not contain gamma-
carboxyglutamic acid.";
RL J. Biol. Chem. 258:5554-5560(1983).
CC -!- FUNCTION: Protein C is a vitamin K-dependent serine protease that
regulates blood coagulation by inactivating factors Va and VIIIa

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CC in the presence of calcium ions and phospholipids.
CC -!- CATALYTIC ACTIVITY: Degradation of blood coagulation factors Va
CC and VIIIa.
CC -!- SUBUNIT: Synthesized as a single chain precursor, which is cleaved
CC into a light chain and a heavy chain held together by a disulfide
CC bond. The enzyme is then activated by thrombin, which cleaves a
CC tetradecapeptide from the amino end of the heavy chain; this is
CC reaction, which occurs at the surface of endothelial cells, is
CC strongly promoted by thrombomodulin.
CC -!- TISSUE SPECIFICITY: Plasma; synthesized in the liver.
CC -!- PM: The vitamin K-dependent, enzymatic carboxylation of some Glu
CC residues allows the modified protein to bind calcium.
CC -!- MISCELLANEOUS: Calcium also binds with stronger affinity to
CC another site, beyond the GLA domain. This GLA-independent binding
CC site is necessary for the recognition of the thrombin-
CC thrombomodulin complex.
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 EGF-like domains.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: K02435; AAA30685.1; -.
CC PIR: A26250; KXBO.
CC HSSP: P04070; IPCU.
CC MEROPS: S01.218; -.
CC InterPro: IPR000152; Asx_hydroxyl_S.
CC InterPro: IPR009003; Cys_Ser_trypsin.
CC InterPro: IPR001881; EGF_Ca.
CC InterPro: IPR006209; EGF_like.
CC InterPro: IPR002383; GLA_blood.
CC InterPro: IPR006210; IEGF.
CC InterPro: IPR001254; Peptidase_S1.
CC InterPro: IPR001314; Peptidase_S1A.
CC InterPro: IPR000294; VitK_dep_GLA.
CC Pfam: PF00008; EGF_2.
CC Pfam: PF00594; gla; 1.
CC Pfam: PF00089; trypsin; 1.
CC PRINTS: PR00722; CHYMOTRYPSIN.
CC PRINTS: PR00001; GLABLOOD.
CC SMART: SM00181; EGF; 2.
CC SMART: SM00069; GLA; 1.
CC SMART: SM00020; TRYD_SPC; 1.
CC PROSITE: PS00010; ASX_HYDROXYL; 1.
CC PROSITE: PS00022; EGF_1; 1.
CC PROSITE: PS01186; EGF_2; 2.
CC PROSITE: PS00026; EGF_3; 1.
CC PROSITE: PS01187; EGF_CA; 1.
CC PROSITE: PS00011; GLU_CARBOXYLATION; 1.
CC PROSITE: PS00240; TRYPSIN_DOM; 1.
CC PROSITE: PS00134; TRYPSIN_HIS; FALSE_NEG.
CC PROSITE: PS00135; TRYPSIN_SER; 1.
CC Blood coagulation; Glycoprotein; Serine protease;
CC Gamma-carboxyglutamic acid; Calcium-binding; Vitamin K; Hydroxylation;
CC EGF-like domain; Repeat; Endothelial cell; Hydrolase; Signal.
CC NON_TER 1 1
CC SIGNAL <1 29
CC PROPEP 30 39
CC CHAIN 40 194
CC CHAIN 197 456
CC PEPTIDE 197 210
CC DOMAIN 94 129
CC DOMAIN 133 173
CC DOMAIN 211 456
CC MOD_RES 45 45
CC MOD_RES 46 46
CC MOD_RES 53 53
CC MOD_RES 55 55
CC PROTEIN C LIGHT CHAIN.
CC PROTEIN C HEAVY CHAIN.
CC ACTIVATION PEPTIDE.
CC EGF-LIKE 1.
CC EGF-LIKE 2.
CC SERINE PROTEASE.
CC GAMMA-CARBOXYGLUTAMIC ACID.
CC GAMMA-CARBOXYGLUTAMIC ACID.
CC GAMMA-CARBOXYGLUTAMIC ACID.

```


like domain in coagulation factor X.";
J. Biol. Chem. 267:19642-19649(1992).
[13]
STRUCTURE BY NMR OF 41-126.
MEDLINE=96387194; PubMed=8794734;
Sunnerhalla M., Olah G.A., Stenflo J., Forsen S., Drakenberg T.,
Sunnerhalla J.;
"The relative orientation of Gla and EGF domains in coagulation
factor X is altered by Ca²⁺ binding to the first EGF domain. A
combined NMR-small angle X-ray scattering study.";
Biochemistry 35:11547-11559(1996)
CC -1- FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that
converts prothrombin to thrombin in the presence of factor Va,
calcium and phospholipid during blood clotting.
CC -1- CATALYTIC ACTIVITY: Preferential cleavage: Arg-|-Thr and then
CC Arg-|-Ile bonds in prothrombin to form thrombin.
CC -1- SUBUNIT: The two chains are formed from a single-chain precursor
by the excision of two Arg residues and are held together by 1 or
more disulfide bonds.
CC -1- PTM: The vitamin K-dependent, enzymatic carboxylation of some
glutamate residues allows the modified protein to bind calcium.
CC -1- PTM: N- and O-glycosylated.
CC -1- PTM: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE
INTRINSIC PATHWAY), OR BY FACTOR VIIA (IN THE EXTRINSIC PATHWAY).
CC -1- MISCELLANEOUS: Calcium also binds, with stronger affinity to
another site, beyond the Gla domain.
CC -1- SIMILARITY: Belongs to peptidase family S1.
CC -1- SIMILARITY: Contains 2 EGF-like domains.
CC
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CC
CC -----
CC EMBL: X00673; CAA25286.1; --
CC PIR: A22867; EXBO.
CC PDB: 1AFO; 31-JAN-94.
CC PDB: 1CCF; 31-MAY-94.
CC PDB: 1WHE; 15-MAY-97.
CC PDB: 1WHF; 15-MAY-97.
CC PDB: 1IOD; 21-JAN-03.
CC PDB: 1KIG; 28-OCT-98.
CC PDB: 1KIG; 28-OCT-98.
CC YEROPS; S01.216; --
CC GlycositesDB; F00743; --
CC InterPro; IPR000152; Asx_hydroxyl_S.
CC InterPro; IPR009003; Cys_Ser_trypsin.
CC InterPro; IPR000742; EGF_2.
CC InterPro; IPR001881; EGF_Ca.
CC InterPro; IPR006209; EGF_like.
CC InterPro; IPR002383; GLA_blood.
CC InterPro; IPR001254; Peptidase_S1.
CC InterPro; IPR001314; Peptidase_S1A.
CC InterPro; IPR000294; VitK_dep_GLA.
CC Pfam; PF00008; EGF_2.
CC Pfam; PF00594; Gla; 1.
CC Pfam; PF00089; trypsin; 1.
CC PRINTS; PR00722; CHYMOTRYPSIN.
CC PRINTS; PR00001; GLABLOOD.
CC SMART; SM00179; EGF_CA; 1.
CC SMART; SM00069; GLA; 1.
CC SMART; SM00020; Tryp_Spc; 1.
CC PROSITE; PS00010; ASX_HYDROXYL; 1.
CC PROSITE; PS00022; EGF_1; 1.
CC PROSITE; PS0186; EGF_2; 2.
CC PROSITE; PS00026; EGF_3; 1.
CC PROSITE; PS01197; EGF_CA; 1.
CC PROSITE; PS00011; GLU CARBOXYLATION; 1.
CC PROSITE; PS00240; TRYPSIN_DOM; 1.
CC PROSITE; PS00134; TRYPSIN_HIS; 1.
CC PROSITE; PS00135; TRYPSIN_SER; 1.

KW Glycoprotein; Hydrolase; Serine protease; Plasma; Blood coagulation;
KW Gamma-carboxyglutamic acid; Hydroxylation; Calcium-binding; Vitamin K;
KW Signal; Zymogen; EGF-like domain; Repeat; Sulfation; 3D-structure.
FT SIGNAL 1 23
FT PROPEP 24 40
FT CHAIN 41 180
FT CHAIN 183 492
FT PROPEP 183 233
FT CHAIN 234 492
FT PROPEP 476 492
FT DOMAIN 86 122
FT DOMAIN 125 165
FT DOMAIN 234 492
FT ACT_SITE 275 275
FT ACT_SITE 321 321
FT ACT_SITE 418 418
FT MOD_RES 46 46
FT MOD_RES 47 47
FT MOD_RES 54 54
FT MOD_RES 56 56
FT MOD_RES 59 59
FT MOD_RES 60 60
Query Match 52.0%; Score 103; DB 1; Length 492;
Best Local Similarity 40.9%; Pred. No. 2.3e-03;
Matches 18; Conservative 8; Mismatches 18; Indels 0; Gaps 0;
QY 1 ANSLFXLRHSSLRXCIXXICDFXKXIFEDVDDTLAFWSKH 44
Db 41 ANSPLEVKQGNLERECLEAEACSLSEAREVFDEAEQDEFWSKY 84
RESULT 8
FA10 HUMAN STANDARD; PRT; 488 AA.
AC P00742; O14340;
DT 21-JUL-1986 (Rel. 01, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Coagulation factor X precursor (EC 3.4.21.6) (Stuart factor).
GN F10.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91216473; PubMed=1902434;
RA Messier T.L., Pittman D.D., Long G.L., Kaufman R.J., Church W.R.;
RT "Cloning and expression in COS-1 cells of a full-length cDNA encoding
RL human coagulation factor X.";
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=87026600; PubMed=3768336;
RA Leytus S.P., Foster D.C., Kurachi K., Davie E.W.;
RT "Gene for human factor X: a blood coagulation factor whose gene
RT organization is essentially identical with that of factor IX and
RL protein C.";
RL Biochemistry 25:5098-5102(1986).
RN [3]
RP SEQUENCE FROM N.A.
RA Rieder M.J., Arnel T.Z., Carrington D.P., Chung M.-W., Lee K.L.,
RA Ozuna M., Peel C.L., Toth E.J., Yi Q., Nickerson D.A.;
RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RC TISSUE=Ovary;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,

RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh P.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonardo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Uedin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S., Lequellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gurnatne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smailus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.,
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences";
RN Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
[5]
RP SEQUENCE OF 13-488 FROM N.A.
RX MEDLINE=85216545; PubMed=2582420;
RA Fung M.R., Hay C.W., McGillivray R.T.A.;
RT "Characterization of an almost full-length cDNA coding for human
RT blood coagulation factor X";
RN Proc. Natl. Acad. Sci. U.S.A. 82:3591-3595(1985).
[6]
RP SEQUENCE OF 19-488 FROM N.A.
RX TISSUE=Liver;
RC MEDLINE=86221713; PubMed=3011603;
RA Kaul R.K., Hildebrand B., Roberts S., Jagadeeswaran P.;
RT "Isolation and characterization of human blood-coagulation factor X
RT cDNA";
RL Gene 41:311-314(1986).
[7]
RP SEQUENCE OF 41-179.
RX MEDLINE=83257207; PubMed=6871167;
RA McMullen B.A., Fujikawa K., Kiesel W., Sasagawa T., Howald W.N.,
RA Kwa E.Y., Weinstein B.;
RT "Complete amino acid sequence of the light chain of human blood
RT coagulation factor X: evidence for identification of residue 63 as
RT beta-hydroxyaspartic acid";
RL Biochemistry 22:2875-2884(1983).
[8]
RP SEQUENCE OF 115-488 FROM N.A., AND TISSUE SPECIFICITY.
RC TISSUE=Liver;
RX MEDLINE=84222026; PubMed=6587384;
RA Leytus S.P., Chung D.W., Kiesel W., Kurachi K., Davie E.W.;
RT "Characterization of a cDNA coding for human factor X";
RL Proc. Natl. Acad. Sci. U.S.A. 81:3699-3702(1984).
[9]
RP SEQUENCE OF 183-234, AND CARBOHYDRATE-LINKAGE SITES.
RX MEDLINE=94062825; PubMed=8243461;
RA Inoue K., Morita T.;
RT "Identification of O-linked oligosaccharide chains in the activation
RT peptides of blood coagulation factor X: the role of the carbohydrate
RT moieties in the activation of factor X";
RL Eur. J. Biochem. 218:153-163(1993).
[10]
RP SEQUENCE OF 1-23 FROM N.A.
RX MEDLINE=90128299; PubMed=2612918;
RA Jagadeeswaran P., Reddy S.V., Rao K.J., Hamsabhusanam K., Lyman G.;
RT "Cloning and characterization of the 5' end (exon 1) of the gene
RT encoding human factor X";
RL Gene 84:517-519(1989).
[11]
RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 86-179 AND 235-278.
RX MEDLINE=93360277; PubMed=8355279;
RA Padmanabhan K., Padmanabhan K.P., Tulinsky A., Park C.H., Bode W.,
RA Huber R., Blankenship D.T., Carlin A.D., Kiesel W.;
RT "Structure of human des(1-45) factor Xa at 2.2-A resolution";
RL J. Mol. Biol. 232:947-966(1993).
[12]
RP X-RAY CRYSTALLOGRAPHY (2.3 ANGSTROMS) OF 86-179 AND 235-278.
RX MEDLINE=98283982; PubMed=9618463;

RA Kamata K., Kawamoto H., Honma T., Iwama T., Kim S.H.;
RT "Structural basis for chemical inhibition of human blood coagulation
RT factor Xa";
RN Proc. Natl. Acad. Sci. U.S.A. 95:6630-6635(1998).
[13]
RP VARIANTS ILE-7 AND HIS-30.
RX MEDLINE=99318093; PubMed=10391209;
RA Carsill M., Altschuler D., Ireland J., Sklar P., Ardlie K., Patil N.,
RA Shaw N., Lane C.R., Lim E.P., Kalyanaraman N., Nemesh J., Ziaugra L.,
RA Friedland L., Rolfe A., Warrington J., Lipshutz R., Daley G.Q.,
RA Lander E.S.;
RT "Characterization of single-nucleotide polymorphisms in coding regions
RT of human genes";
RN Nat. Genet. 22:231-238(1999).
[14]
RP ERRATUM.
RA Carsill M., Altschuler D., Ireland J., Sklar P., Ardlie K., Patil N.,
RA Shaw N., Lane C.R., Lim E.P., Kalyanaraman N., Nemesh J., Ziaugra L.,
RA Friedland L., Rolfe A., Warrington J., Lipshutz R., Daley G.Q.,
RA Lander E.S.;
RT "FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that
RT converts prothrombin to thrombin in the presence of factor Va,
RT calcium and phospholipid during blood clotting."
CC -! CATALYTIC ACTIVITY: Preferential cleavage: Arg-|-Thr and then
CC Arg-|-Ile bonds in prothrombin to form thrombin.
CC -! SUBUNIT: The two chains are formed from a single-chain precursor
CC by the excision of two Arg residues and are held together by 1 or
CC more disulfide bonds.
CC -! TISSUE SPECIFICITY: Plasma; synthesized in the liver.
CC -! PTM: The vitamin K-dependent, enzymatic carboxylation of some
CC glutamate residues allows the modified protein to bind calcium.
CC -! PTM: N- and O-glycosylated.
CC -! PTM: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE
CC INTRINSIC PATHWAY), OR BY FACTOR VIIA (IN THE EXTRINSIC PATHWAY).
CC -! SIMILARITY: Belongs to peptidase family S1.
CC -! SIMILARITY: Contains 2 EGF-like domains.

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CC or send an email to license@isb-sib.ch).

EMBL; K03194; AAA52490.1; -
EMBL; M57285; AAA52421.1; -
EMBL; AF503510; AAM19347.1; -
EMBL; BC046125; AAH46125.1; -
EMBL; L29433; AAA52764.1; -
EMBL; L29433; AAA52764.1; JOINED.
EMBL; L00390; AAA52764.1; JOINED.
EMBL; L00391; AAA52764.1; JOINED.
EMBL; L00392; AAA52764.1; JOINED.
EMBL; L00393; AAA52764.1; JOINED.
EMBL; L00394; AAA52764.1; JOINED.
EMBL; L00395; AAA52764.1; JOINED.
EMBL; L00396; AAA52764.1; JOINED.
EMBL; M22613; AAA51984.1; -
EMBL; K01886; AAA52486.1; -
EMBL; M33297; AAA52636.1; -
PIR; A24478; EXHU.
PDB; 1HCG; 08-MAY-95.
PDB; 1FAX; 29-OCT-97.
PDB; 1FXV; 17-JUN-98.
PDB; 1XKA; 23-MAR-99.
PDB; 1XKB; 23-MAR-99.
PDB; 1EZQ; 20-SEP-00.
PDB; 1FOR; 20-SEP-00.
PDB; 1FOS; 20-SEP-00.
PDB; 1FJS; 17-NOV-00.
PDB; 1G2L; 20-OCT-01.
PDB; 1G2M; 20-OCT-01.


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FT DOMAIN          20      78    EXTRACELLULAR (POTENTIAL).
FT TRANSMEM       79     101    POTENTIAL.
FT DOMAIN        102     231    CYTOPLASMIC (POTENTIAL).
FT DOMAIN         23   60      GLA-RICH.
SQ SEQUENCE      231 AA; 25948 MW; 8A373E484949D81 CRC64;

Query Match           48.5%; Score 96; DB 1; Length 231;
Best Local Similarity 39.0%; Pred. No. 1.4e-08;
Matches 16; Conservative 8; Mismatches 17; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLXRXCIXXICDFXXAKXIPEVDVDTLAFW 41
   |||||:::||:::||::||:||||:|
DB 20 ANEFLEELRQTIERCEMEIECSYEVEKVENKEKTMEFW 60

RESULT 10
FA10_RABIT STANDARD; PRT; 490 AA.
ID AC Q19045;
DT DT 15-DEC-1998 (Rel. 37, Created)
DI DI 15-DEC-1998 (Rel. 37, Last sequence update)
DE DE 10-OCT-2003 (Rel. 42, Last annotation update)
DN DN Coagulation factor X precursor (EC 3.4.21.6) (Stuart factor).
GN GN F10.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxId=9986;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97256311; PubMed=9101642;
RA Pandurthi U.R.; Anderson K.D.; James H.L.;
RT "Characterization of a full-length cDNA for rabbit factor X.";
RL Thromb. Res. 85:503-514 (1997).
CC -|- FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that
CC converts prothrombin to thrombin in the presence of factor Va,
CC calcium and phospholipid during blood clotting.
CC -|- CATALYTIC ACTIVITY: Preferential cleavage: Arg--|Thr and then
CC Arg--|Ile bonds in prothrombin to form thrombin.
CC -|- SUBUNIT: The two chains are formed from a single-chain precursor
CC by the excision of two arg residues and are held together by 1 or
CC more disulfide bonds.
CC -|- PTM: The vitamin K-dependent, enzymatic carboxylation of some
CC glutamate residues allows the modified protein to bind calcium (By
CC similarity).
CC -|- PTM: N- and O-glycosylated (By similarity).
CC -|- PTM: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE
CC INTRINSIC PATHWAY), OR BY FACTOR VIIA (IN THE EXTRINSIC PATHWAY)
CC (BY SIMILARITY).
CC -|- MISCELLANEOUS: Calcium also binds, with stronger affinity to
CC another site, beyond the GLA domain.
CC -|- SIMILARITY: Belongs to peptidase family S1.
CC -|- SIMILARITY: Contains 2 EGF-like domains.
-----
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CC or send an email to licens@isb-sib.ch).
-----
CC EMBL; AF003200; AAB62542.1; -.
CC HSP; P00742; IHCG.
DR MEROPS; S01_216; -.
DR InterPro; IPR000152; Asx_hydroxyl_S.
DR InterPro; IPR009003; Cys_Ser_trypsin.
DR InterPro; IPR000742; EGF 2.
DR InterPro; IPR001881; EGF Ca.
DR InterPro; IPR001438; EGF II.
DR InterPro; IPR006209; EGF_like.
DR InterPro; IPR002383; GLA_blood.

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DR InterPro: IPR001438; EGF II.
DR InterPro: IPR006209; EGF-like.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Peptidase_S1.
DR InterPro: IPR001314; Peptidase_S1A.
DR InterPro: IPR000294; VitK_dep_GLA.
DR Pfam: PF00008; EGF; 2.
DR Pfam: PF00594; gla; 1.
DR Pfam: PF00089; trypsin; 1.
DR PRINTS: PR00722; CHYMOTRYPSIN.
DR PRINTS: PR00010; EGF_blood.
DR PRINTS: PR00001; GLABLOOD.
DR SMART: SM00179; EGF_CA; 1.
DR SMART: SM00069; GLA; 1.
DR SMART: SM00020; Tryp SPC; 1.
DR PROSITE: PS00010; ASX HYDROXYL; 1.
DR PROSITE: PS00022; EGF_1; 1.
DR PROSITE: PS01186; EGF_2; 1.
DR PROSITE: PS00026; EGF_3; 1.
DR PROSITE: PS01187; EGF_CA; 1.
DR PROSITE: PS00011; GLU CARBOXYLATION; 1.
DR PROSITE: PS50240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
DR Hydrolase; Serine protease; Blood coagulation; Zymogen; Glycoprotein;
KW Liver; Plasma; Vitamin K; Calcium-binding; Gamma-carboxyglutamic acid;
KW EGF-like domain; Repeat; Signal; Hydroxylation.
FT SIGNAL 1 21
FT PROPEP 22 39
FT CHAIN 40 191
FT CHAIN 192 444
FT DOMAIN 45 74
FT DOMAIN 85 121
FT DOMAIN 126 167
FT DOMAIN 192 444
FT SITE 191 192
FT ACT_SITE 232 232
FT ACT_SITE 281 281
FT ACT_SITE 383 383
FT BINDING 377 377
FT DISULFID 56 61
FT DISULFID 89 100
FT DISULFID 94 109
FT DISULFID 111 120
FT DISULFID 130 141
FT DISULFID 137 151
FT DISULFID 153 166
FT DISULFID 174 301
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FT MOD_RES 64 64
FT MOD_RES 65 65
FT MOD_RES 68 68
FT MOD_RES 74 74
FT MOD_RES 102 102
FT CARBOHYD 211 211
FT CARBOHYD 242 242
FT CARBOHYD 306 306
SQ SEQUENCE 444 AA; 49011 MW; 0481ABC4FE5427F8 CRC64;

Query Match

Best Local Similarity 47.5%; Score 94; DB 1; Length 444;

Matches 18; Conservative 5; Mismatches 16; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSLRXCIIXICDFXXAKXIPEDVDDTLAFW 41
DB 40 ANSFLLELRPGSLRECKELCSFEAREVPQSTERTKQFW 80
RESULT 12
FA7_HUMAN
ID FA7_HUMAN STANDARD; PRT; 466 AA.
AC P08709; Q14339;
DT 01-JAN-1988 (Rel. 06, Created)
DT 01-JAN-1988 (Rel. 06, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Coagulation factor VII precursor (EC 3.4.21.2) (Serum prothrombin
conversion accelerator) (Bptacog alfa).
GN F7.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RC TISSUE=Liver;
RX MEDLINE=86205965; PubMed=3486420;
RA Hagen F.S., Gray C.L., O'Hara P.J., Grant F.J., Saari G.C.,
RA Woodbury R.G., Hart C.E., Inley M.Y., Kisiel W., Kurachi K.,
RA Davie E.W.;
RT "Characterization of a cDNA coding for human factor VII.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:2412-2416(1986).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=87260948; PubMed=3037537;
RA O'Hara P.J., Grant F.J., Haldeman B.A., Gray C.L., Inley M.Y.,
RA Hagen F.S., Murray M.J.;
RT "Nucleotide sequence of the gene coding for human factor VII, a
vitamin K-dependent protein participating in blood coagulation.";
RL Proc. Natl. Acad. Sci. U.S.A. 84:5158-5162(1987).
RN [3]
RP SEQUENCE FROM N.A., AND VARIANTS THR-352; GLN-413 AND LYS-445.
RA Rieder M.J., Armet T.Z., Carrington D.P., Chung M.-W., Lee K.L.,
RA Poel C.L., Toth E.J., Yi Q., Nickerson D.A.;
RL Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE OF 61-466, AND POST-TRANSLATIONAL MODIFICATIONS.
RX MEDLINE=8908153; PubMed=3264725;
RA Thim L., Bjoern S., Christensen M., Nicolaisen E.M., Lund-Hansen T.,
RA Pedersen A.H., Hedner U.;
RT "Amino acid sequence and posttranslational modifications of human
factor VIIa from plasma and transfected baby hamster kidney cells.";
RL Biochemistry 27:7785-7793(1988).
RN [5]
RP CARBOHYDRATE-LINKAGE SITES SER-112 AND SER-120.
RX MEDLINE=91250411; PubMed=1904059;
RA Bjoern S., Foster D.C., Thim L., Wibeberg F.C., Christensen M.,
RA Komiyama Y., Pedersen A.H., Kisiel W.;
RT "Human plasma and recombinant factor VII. Characterization of O-
glycosylations at serine residues 52 and 60 and effects of site-
directed mutagenesis of serine 52 to alanine.";
RL J. Biol. Chem. 266:11051-11057(1991).
RN [6]
RP STRUCTURE OF CARBOHYDRATE ON SER-112.
RX MEDLINE=90062160; PubMed=2511201;
RA Nishimura H., Kawabata S., Kisiel W., Hase S., Ikenaka T., Takao T.,
RA Shimonishi Y., Iwanaga S.;
RT "Identification of a disaccharide (Xyl-Glc) and a trisaccharide
(Xyl2-Glc) O-glycosidically linked to a serine residue in the first
epidermal growth factor-like domain of human factors VII and IX and
protein Z and bovine protein Z.";
RL J. Biol. Chem. 264:20320-20325(1989).
RN [7]
RP STRUCTURE OF CARBOHYDRATE ON SER-112.
RX MEDLINE=91344709; PubMed=219367;
RA Iwanaga S., Nishimura H., Kawabata S., Kisiel W., Hase S., Ikenaka T.;
RT "A new trisaccharide sugar chain linked to a serine residue in the

first EGF-like domain of clotting factors VII and IX and protein Z.",
Adv. Exp. Med. Biol. 281:121-131(1990).
[8]
X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS) OF FVIIA IN COMPLEX WITH TF.
MEDLINE=96175641; PubMed=8598903;
Banner D.W., D'Arcy A., Chene C., Winkler F.K., Guha A.,
Königsberg W.H., Nemtsov Y., Kirchofer D.;
The crystal structure of the human coagulation factor
VIIa with soluble tissue factor.";
Nature 380:41-46(1996).
[9]
X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF FVIIA IN COMPLEX WITH TF.
MEDLINE=99126538; PubMed=9925787;
Zhang E., St Charles R., Tulinsky A.;
"Structure of extracellular tissue factor complexed with factor VIIa
inhibited with a BPTI mutant.";
J. Mol. Biol. 285:2089-2104(1999).
[10]
STRUCTURE BY NMR OF 105-145.
MEDLINE=98367502; PubMed=9692950;
Muranyi A., Finn B.E., Gippert G.P., Forsen S., Stenflo J.,
Drakenberg T.;
"Solution structure of the N-terminal EGF-like domain from human
factor VII.";
Biochemistry 37:10605-10615(1998).
[11]
VARIANT GLN-364.
MEDLINE=91300046; PubMed=2070047;
O'Brien D.P., Gale K.W., Anderson J.S., McVey J.H., Miller G.J.,
Meade T.W., Tuddenham E.G.D.;
"Purification and characterization of factor VII 304-Gln: a variant
molecule with reduced activity isolated from a clinically unaffected
male.";
Blood 78:132-140(1991).
[12]
VARIANTS GLN-364 AND PHE-370.
MEDLINE=92340074; PubMed=1634227;
Marchetti G., Patraccchini P., Gemmati D., Derosa V., Pinotti M.,
Rodorigo G., Casonato A., Girolami A., Bernardi F.;
"Detection of two missense mutations and characterization of a repeat
polymorphism in the factor VII gene (F7).";
Hum. Genet. 89:497-502(1992).
[13]
VARIANT TYR-239.
MEDLINE=93372811; PubMed=8364544;
Marchetti G., Ferrati M., Patraccchini P., Redaelli R., Bernardi F.;
"A missense mutation (178Cys--Tyr) and two neutral dimorphisms
(115His and 338Ser) in the human coagulation factor VII gene.";
Hum. Mol. Genet. 2:1055-1056(1993).
[14]
VARIANTS.
MEDLINE=94061028; PubMed=8242057;
Takamiya O., Kemball-Cook G., Martin D.M.A., Cooper D.N.,
Tamura A., Meili E., Hahn I., Prangnell D.R., Lumley H.,
Tuddenham E.G.D., McVey J.H.;
"Detection of missense mutations by single-strand conformational
polymorphism (SSCP) analysis in five dysfunctional variants of
coagulation factor VII.";
Hum. Mol. Genet. 2:1355-1359(1993).
[15]
VARIANTS CHARLOTTE GLN-139 AND GLN-212.
MEDLINE=94264305; PubMed=8204879;
Chang S., Clarke B., Sridhara S., Chu K., Friedman P., Vandusen W.,
Roberts H.R., Blajchman M., Monroe D.M., High K.A.;
"Severe factor VII deficiency caused by mutations abolishing the
cleavage site for activation and altering binding to tissue factor.";
Blood 83:3524-3535(1994).
[16]
VARIANT VAL-354.
MEDLINE=95072589; PubMed=7991691;
Bernardi F., Castaman G., Redaelli R., Pinotti M., Lunghi B.,
Rodeghiero F., Marchetti G.;
"Topologically equivalent mutations causing dysfunctional coagulation

factors VII (294Ala-->Val) and X (334Ser-->Pro).";
 Hum. Mol. Genet. 3:1175-1177(1994).
 [17]
 VARIANT MIE HIS-307.
 MEDLINE=95064562; PubMed=7974346;
 Ohiva W., Hayashi T., Wada H., Minamikawa K., Shirakawa S.,
 Suzuki K.;
 "Factor VII MIE: homozygous asymptomatic type I deficiency caused by
 an amino acid substitution of His (CAC) for Arg(247) (CGC) in the
 catalytic domain.";
 Thromb. Haemost. 71:773-777(1994).
 [18]
 VARIANT MET-419.
 MEDLINE=96247510; PubMed=8652821;
 Arbini A.A., Mannucci P.M., Bauer K.A.;
 "A thr359Met mutation in factor VII of a patient with a hereditary
 deficiency causes defective secretion of the molecule.";
 Blood 87:5085-5094(1996).
 [19]
 VARIANTs TRP-283; LYS-325; VAL-358; GLN-364; GLU-402 AND GLN-413.
 MEDLINE=97001216; PubMed=8844208;
 Bernardi F., Castaman G., Pinotti M., Ferraresi P., di Iasio M.G.,
 Lunghi B., Rodeghiero F., Marchetti G.;
 "Mutation pattern in clinically asymptomatic coagulation factor VII
 deficiency.";
 Hum. Mutat. 8:108-115(1996).
 [20]
 VARIANT VAL-304.
 MEDLINE=97037613; PubMed=8883260;
 Tamary H., Fromovich Y., Shamlon L., Reich Z., Dym O., Lanir N.,
 Brenner B., Paz M., Luder A.S., Blau O., Korostishevsky M.,
 Zaitov R., Seligsohn U.;
 "A424Val is a common, probably ancient mutation causing factor VII
 deficiency in Moroccan and Iranian Jews.";
 Thromb. Haemost. 76:283-291(1996).
 [21]
 VARIANT MORIOKA PRO-13.
 MEDLINE=98235713; PubMed=9576180;
 Ozawa T., Takikawa Y., Niya K., Ejiri N., Suzuki K., Sato S.,
 Sakuragawa N.;
 "Factor VII Moriooka (FVII L-26P): a homozygous missense mutation in
 the signal sequence identified in a patient with factor VII
 deficiency.";
 Br. J. Haematol. 101:47-49(1998).
 [22]
 VARIANTS MALTA THR-194 AND VAL-304.
 MEDLINE=98112461; PubMed=9452082;
 Alehinawi C., Scerri C., Galdies R., Aquilina A., Felice A.E.;
 "Two new missense mutations (P14F and A244V) in the coagulation
 factor VII gene.";
 Hum. Mutat. Suppl. 1:S189-S191(1998).
 [23]
 VARIANTS ASP-295 AND GLN-413.
 MEDLINE=99318093; PubMed=10391209;
 Cartgill M., Althahuler D., Ireland J., Sklar P., Ardlie K., Patil N.,
 Shaw N., Lane C.R., Lim E.P., Kalyanaraman N., Nimesh J., Ziaugra L.,
 Friedland J., Rolfe A., Warrington J., Lipschutz R., Daley G.Q.,
 Lander E.S.;
 "Characterization of single-nucleotide polymorphisms in coding regions
 of human genes.";
 Nat. Genet. 22:231-238(1999).
 Query Match 46.5%; Score 92; DB 1; Length 466;
 Best Local Similarity 46.3%; Pred. No. 1 Se-07;
 Matches 19; Conservative 4; Mismatches 18; Indels 0; Gaps
 1 ANSFLXRLRSLRXICIXICDFXKXKIFEDVDOTLAFW 41
 61 ANAFLEIRPGSLERECKQCSFEAREIFKDAERTKLFW 101

RESULT 13
TMG1 HUMAN

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ID TMG1_HUMAN STANDARD; PRT; 218 AA.
AC O14658;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Transmembrane gamma-carboxyglutamic acid proprotein (Proline-
DE rich gla protein 1) (Proline-rich gamma-carboxyglutamic acid protein
DE 1).
GN PRRG1 OR TMG1 OR PRGP1..
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP MEDLINE=97404347; PubMed=9256434;
RX Kulkarni J.D., Harris J.E., Haldeman B.A., Davie B.W.;
RA "Primary structure and tissue distribution of two novel proline-rich
RT gamma-carboxyglutamic acid proteins.";
RL Proc. Natl. Acad. Sci. U.S.A. 94:9058-9062(1997).
CC -!- TISSUE SPECIFICITY: Highly expressed in the spinal cord.
CC -!- PTM: Gla residues are produced after subsequent posttranslational
CC modifications of glutamate by a vitamin K-dependent gamma-
CC carboxylase.
CC
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CC
CC -----
CC EMBL; AF009242; AAB67070.1; -.
CC HSP; P00740; 1CFH.
CC Genew; HGNC:9469; -PRRG1.
CC MIM; 604428; -.
CC GO; GO:0005887; C:integral to plasma membrane; TAS.
CC InterPro; IPR002383; GLA_blood.
CC InterPro; IPR000294; VitK_dep_GLA.
CC Pfam; PF00594; gla; 1.
CC PRINTS; PR00001; GLABLOOD.
CC SMART; SM00069; GLA; 1.
CC PROSITE; PS00011; GLU CARBOXYLATION; 1.
CC Gamma-carboxyglutamic acid; Vitamin K; Transmembrane.
KW PROPEP
FT CHAIN 1 20
FT TRANSMEMBRANE GAMMA-CARBOXYGLUTAMIC ACID
FT PROTEIN 1.
FT DOMAIN 21 83
FT TRANSMEM 84 106
FT DOMAIN 107 218
FT CYTOPLASMIC (POTENTIAL).
FT DOMAIN 24 61
FT GLA-RICH.
FT DOMAIN 131 135
FT POLY-PRO.
SQ SEQUENCE 218 AA; 24947 MW; 26538A61AB0AE98 CRC64;

Query Match
Best Local Similarity 34.1%; Score 81; DB 1; Length 218;
Matches 15; Conservative 7; Mismatches 22; Indels 0; Gaps 0;

QY 1 ANSFLXLIHSHSLXKXCIXICDFXXKXIFEDVDDTLAFWSKH 44
DB 21 ANGFFFEIRQGNIECKEEFCTFEAREAFENNEKTEFWSTY 64

RESULT 14
THRB_RAT
ID THRB_RAT STANDARD; PRT; 617 AA.
AC P18292;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Prothrombin precursor (EC 3.4.21.5).
DE F2.
GN
```

```
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Sprague-Dawley; TISSUE=Liver;
RX MEDLINE=90332426; PubMed=2377469;
RA Dhanich M., Monard D.;
RT "cDNA sequence of rat prothrombin.";
RL Nucleic Acids Res. 18:4251-4251(1990).
RN [2]
RP SEQUENCE OF 383-617 FROM N.A.
RC TISSUE=Liver;
RX MEDLINE=92212913; PubMed=1557383;
RA Banfield D.K., Macgillivray R.T.;
RT "Partial characterization of vertebrate prothrombin cDNAs:
RT amplification and sequence analysis of the B chain of thrombin from
RT nine different species.";
RL Proc. Natl. Acad. Sci. U.S.A. 89:2779-2783(1992).
CC -!- FUNCTION: THROMBIN, WHICH CLEAVES BONDS AFTER ARG & LYS, CONVERTS
CC FIBRINOGEN TO FIBRIN AND ACTIVATES FACTORS V, VII, VIII, XIII,
CC AND, IN COMPLEX WITH THROMBOMODULIN, PROTEIN C.
CC -!- CATALYTIC ACTIVITY: Preferential cleavage: Arg-Gly; activates
CC fibrinogen to fibrin and releases fibrinopeptide A and B.
CC -!- PTM: THE GAMMA-CARBOXYGLUTAMYL RESIDUES, WHICH BIND CALCIUM IONS,
CC RESULT FROM THE CARBOXYLATION OF GLUTAMYL RESIDUES BY A MICROSOMAL
CC ENZYME, THE VITAMIN K-DEPENDENT CARBOXYLASE. THE MODIFIED RESIDUES
CC ARE NECESSARY FOR THE CA-DEPENDENT INTERACTION WITH A NEGATIVELY
CC CHARGED PHOSPHOLIPID SURFACE, WHICH IS ESSENTIAL FOR THE CONVERSION
CC OF PROTHROMBIN TO THROMBIN.
CC -!- MISCELLANEOUS: PROTHROMBIN IS ACTIVATED ON THE SURFACE OF A
CC PHOSPHOLIPID MEMBRANE THAT BINDS THE AMINO END OF PROTHROMBIN &
CC FACTORS VA & XA IN CA-DEPENDENT INTERACTIONS. FACTOR XA REMOVES
CC THE ACTIVATION PEPTIDE & CLEAVES THE REMAINING PART INTO LIGHT &
CC HEAVY CHAINS. THE ACTIVATION PROCESS STARTS SLOWLY BECAUSE FACTOR
CC V ITSELF HAS TO BE ACTIVATED BY THE INITIAL, SMALL AMOUNTS OF
CC THROMBIN.
CC -!- MISCELLANEOUS: Thrombin can itself cleave the amino terminal
CC fragment (fragment 1) of the prothrombin, prior to its activation
CC by factor Xa.
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 kringle domains.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC use by non-profit institutions as long as its content is in no way
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CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC -----
CC EMBL; X52835; CAA37017.1; -.
CC EMBL; M81397; AAA42240.1; -.
CC PIR; S10511; S10511.
CC HSP; P00734; IUVS.
CC MEQPS; S01.217; -.
CC InterPro; IPR009003; Cys_Ser_trypsin.
CC InterPro; IPR002383; GLA_blood.
CC InterPro; IPR000001; Kringle.
CC InterPro; IPR001254; Peptidase_S1.
CC InterPro; IPR001314; Peptidase_S1A.
CC InterPro; IPR003966; Peptidase_S1A_pr.
CC InterPro; IPR000294; VitK_dep_GLA.
CC Pfam; PF00594; gla; 1.
CC Pfam; PF00051; kringle; 2.
CC Pfam; PF00089; trypsin; 1.
CC PRINTS; PR00722; CHYMOTRYPSIN.
CC PRINTS; PR00001; GLABLOOD.
CC PRINTS; PR00018; KRINGLE.
CC PRINTS; PR01505; PROTHROMBIN.
CC ProDom; PD000395; Kringle; 2.
CC SMART; SM00069; GLA; 1.
```

DR SMART; SMO0130; KR; 2.
DR PROSITE; SMO0020; TRYP_SPC; 1.
DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
DR PROSITE; PS00021; KRINGLE_1; 2.
DR PROSITE; PS00021; KRINGLE_2; 2.
DR PROSITE; PS00070; KRINGLE_2; 2.
DR PROSITE; PS00240; TRYP_SIN_DOM; 1.
DR PROSITE; PS00134; TRYP_SIN_DOM; 1.
DR PROSITE; PS00135; TRYP_SIN_SER; 1.
KW Blood coagulation; Plasma; Calcium-binding; Glycoprotein; Repeat;
KW Vitamin K; Zymogen; Gamma-carboxyglutamic acid; Acute phase; Liver;
KW Hydroxylase; Serine protease; Kringle; Signal.
FT SIGNAL 1 24
FT PROPEP 25 43
FT CHAIN 44 617
FT PEPTIDE 44 200
FT PEPTIDE 201 323
FT PEPTIDE 324 359
FT CHAIN 360 617
FT CHAIN 109 187
FT DOMAIN 215 292
FT DOMAIN 360 617
FT SITE 200 201
FT SITE 323 324
FT SITE 359 360
FT SITE 402 402
FT ACT_SITE 458 458
FT ACT_SITE 564 564
FT MOD_RES 50 50
FT MOD_RES 51 51
FT MOD_RES 58 58
FT MOD_RES 60 60
FT MOD_RES 63 63
FT MOD_RES 64 64
FT MOD_RES 69 69
FT MOD_RES 70 70
FT MOD_RES 73 73
FT MOD_RES 76 76
FT CARBOHYD 120 120
FT CARBOHYD 144 144
FT CARBOHYD 412 412
FT CARBOHYD 552 552
FT CARBOHYD 61 66
FT DISULFID 91 104
FT DISULFID 109 187
FT DISULFID 130 170
FT DISULFID 158 182
FT DISULFID 215 292
FT DISULFID 236 276
FT DISULFID 264 287
FT DISULFID 332 478
FT DISULFID 387 403
FT DISULFID 532 546
FT DISULFID 560 590
SQ SEQUENCE 617 AA; AD27D1B71445DB1D CRC64;
Query Match 39.6%; Score 78.5; DB 1; Length 617;
Best Local Similarity 40.0%; Pred. No. 3.7e-05;
Matches 18; Conservative 5; Mismatches 21; Indels 1; Gaps 1;
QY 1 ANS-FLXLRHSLSXRCIXXICDPXXAXXIFEDVDTLAFSKH 44
DB 44 ANSGFLELRKGNLEECVCEQCSYBEAFEALESPOQTDVFWAKY 88
RESULT 15
THRB_MOUSE
ID THRB_MOUSE STANDARD; PRT; 618 AA.
AC P19221;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Prothrombin precursor (EC 3.4.21.5).
GN F2 OR CF2.

OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6; TISSUE=Liver;
RX MEDLINE=91025551; PubMed=2222810;
RA Fritzen Degen S.J., Schaffer L.A., Jamison C.S., Grant S.G.,
RA Fitzgibbon J.J., Pai J.-A., Chapman V.M., Elliott R.W.;
RT "Characterization of the cDNA coding for mouse prothrombin and
RT localization of the gene on mouse chromosome 2.";
RL DNA Cell Biol. 9:487-498(1990).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=FVB/N; TISSUE=Liver;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.B.,
RA Brownstein M.J., Ustin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Locuillano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettner M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalls D.B.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [3]
RP SEQUENCE OF 394-618 FROM N.A.
RC TISSUE=Liver;
RX MEDLINE=92212913; PubMed=1557383;
RA Banfield D.K., Macgillivray R.T.;
RT "Partial characterization of vertebrate prothrombin cDNAs:
RT amplification and sequence analysis of the B chain of thrombin from
RT nine different species";
RL Proc. Natl. Acad. Sci. U.S.A. 89:2779-2783(1992).
CC -!- FUNCTION: Thrombin, which cleaves bonds after Arg and Lys,
CC converts fibrinogen to fibrin and activates factors V, VII, VIII,
CC XIII, and, in complex with thrombomodulin, protein C.
CC -!- CATALYTIC ACTIVITY: Preferential cleavage: Arg-Gly; activates
CC fibrinogen to fibrin and releases fibrinopeptide A and B.
CC -!- PTM: THE GAMMA-CARBOXYGLUTAMYL RESIDUES, WHICH BIND CALCIUM IONS,
CC RESULT FROM THE CARBOXYLATION OF GLUTAMYL RESIDUES BY A MICROSOMAL
CC ENZYME, THE VITAMIN K-DEPENDENT CARBOXYLASE. THE MODIFIED RESIDUES
CC ARE NECESSARY FOR THE CA-DEPENDENT INTERACTION WITH A NEGATIVELY
CC CHARGED PHOSPHOLIPID SURFACE, WHICH IS ESSENTIAL FOR THE CONVERSION
CC OF PROTHROMBIN TO THROMBIN.
CC -!- MISCELLANEOUS: Prothrombin is activated on the surface of a
CC phospholipid membrane that binds the amino end of prothrombin and
CC factors Va and Xa in Ca-dependent interactions; factor Xa removes
CC the activation peptide and cleaves the remaining part into light
CC and heavy chains. The activation process starts slowly because
CC factor V itself has to be activated by the initial, small amounts
CC of thrombin.
CC -!- MISCELLANEOUS: Thrombin can itself cleave the amino terminal
CC fragment (fragment 1) of the prothrombin, prior to its activation
CC by factor Xa.
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 kringle domains.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its

Job time : 11.5 secs

APR 17 1964

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OM protein - protein search, using sw model

Run on: March 1, 2004, 09:55:12 ; Search time 37.5 Seconds
(without alignments)
370.208 Million cell updates/sec

Title: SEQ1-32GLU-33ASP

Perfect score: 198

Sequence: 1 ANSFLXLRHSLRXCIXX.....XXAKXIFedVDDTLAFWSKH 44

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTRMBL.25.*

1: sp_archaea.*

2: sp_bacteria.*

3: sp_fungi.*

4: sp_human.*

5: sp_invertebrate.*

6: sp_mammal.*

7: sp_mhc.*

8: sp_organelle.*

9: sp_phage.*

10: sp_plant.*

11: sp_rodent.*

12: sp_virus.*

13: sp_vertebrate.*

14: sp_unclassified.*

15: sp_virus.*

16: sp_bacteriap.*

17: sp_archaeap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	144	72.7	456	6 Q9TTR0	Q9TTR0 canis famil
2	133	67.2	460	11 Q31WN8	Q31WN8 mus musculu
3	130	65.7	55	4 Q8J002	Q8J002 homo sapien
4	130	65.7	55	4 Q8IXB5	Q8IXB5 homo sapien
5	127	64.1	460	11 Q99PC6	Q99PC6 mus musculu
6	107	54.0	482	11 Q63207	Q63207 rattus norv
7	102	51.5	455	13 Q7SY86	Q7SY86 xenopus lae
8	100	50.5	433	13 Q804X6	Q804X6 gallus gall
9	98	48.7	524	13 Q7SXH8	Q7SXH8 brachydanio
10	96.5	48.7	443	13 Q8JHC9	Q8JHC9 brachydanio
11	96	48.5	231	4 Q8N2N6	Q8N2N6 homo sapien
12	93	47.0	340	11 Q80Y26	Q80Y26 mus musculu
13	93	47.0	434	13 Q7T3B6	Q7T3B6 brachydanio
14	93	47.0	481	11 Q54740	Q54740 mus musculu
15	93	47.0	481	11 Q99L32	Q99L32 mus musculu
16	93	47.0	481	11 Q88947	Q88947 mus musculu

17	92	46.5	679	4 Q86PQ8	Q86PQ8 homo sapien
18	89	44.9	474	13 Q8JHC8	Q8JHC8 brachydanio
19	88.5	44.7	442	13 Q804X1	Q804X1 fugu rubrip
20	87	43.9	469	6 Q9GMD9	Q9GMD9 ornithorhyn
21	82	41.4	229	13 Q8JU40	Q8JU40 xenopus lae
22	81	40.9	268	4 Q8NEK6	Q8NEK6 homo sapien
23	80	40.4	425	13 Q804X7	Q804X7 gallus gall
24	80	40.4	612	13 Q804W7	Q804W7 fugu rubrip
25	79	39.9	446	11 Q8K3U6	Q8K3U6 rattus norv
26	76	38.4	100	4 Q15253	Q15253 homo sapien
27	76	38.4	622	4 Q7Z7P3	Q7Z7P3 homo sapien
28	74	37.4	471	13 Q804X6	Q804X6 gallus gall
29	74	37.4	475	13 Q804W9	Q804W9 fugu rubrip
30	74	37.4	497	4 Q7Z7I5	Q7Z7I5 homo sapien
31	74	37.4	650	4 Q165I9	Q165I9 homo sapien
32	74	37.4	650	4 Q9NSD0	Q9NSD0 homo sapien
33	73	36.9	446	11 Q61109	Q61109 mus musculu
34	71.5	36.1	542	5 Q8T6I3	Q8T6I3 halocynthia
35	71	35.9	138	6 Q28994	Q28994 sus scrofa
36	71	35.9	441	13 Q804X2	Q804X2 fugu rubrip
37	70	35.4	607	13 Q91001	Q91001 gallus gall
38	69	34.8	461	6 Q95ND7	Q95ND7 pan troglod
39	69	34.8	461	6 Q95ND6	Q95ND6 pan troglod
40	67	33.8	648	6 Q29094	Q29094 sus scrofa
41	66.5	33.6	433	13 Q8JHD0	Q8JHD0 brachydanio
42	66.5	33.6	433	13 Q90VK1	Q90VK1 brachydanio
43	66	33.3	49	6 Q95ME8	Q95ME8 bos taurus
44	66	33.3	52	4 Q8IXD5	Q8IXD5 homo sapien
45	64	32.3	430	13 Q804X0	Q804X0 fugu rubrip

ALIGNMENTS

RESULT 1

ID	Q9TTR0	PRELIMINARY;	PRT;	456 AA.
AC	Q9TTR0;			
DT	01-MAY-2000 (TrEMBLrel. 13, Created)			
DT	01-MAY-2000 (TrEMBLrel. 13, Last sequence update)			
DT	01-OCT-2003 (TrEMBLrel. 25, Last annotation update)			
DS	Protein C precursor.			
GN	PROC.			
OS	Canis familiaris (Dog).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.			
OX	NCBI_TaxID=9615;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RA	Leeb T., Kopp T., Deppe A., Breen M., Matis U., Brunnberg L.,			
RA	Brenig B.;			
RT	"Molecular characterization and chromosomal assignment of the canine			
RT	protein C gene."			
RL	Mamm. Genome 10:135-139(1999).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
PX	MEDLINE=9371952; Pubmed=10443005;			
RA	Leeb T., Pfeiffer I., Kopp T., Deppe A., Brenig B.;			
RT	"Analysis of canine protein C gene polymorphisms."			
RL	Anim. Genet. 30:237-238(1999).			
CC	-!- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.			
DR	EMBL; AJ001979; CAA05126.1; -			
DR	HSP; P04070; IAUU			
DR	GO; GO:0005576; C:extracellular; IEA.			
DR	GO; GO:0005509; F:calcium ion binding; IEA.			
DR	GO; GO:0004263; F:chymotrypsin activity; IEA.			
DR	GO; GO:0008233; F:peptidase activity; IEA.			
DR	GO; GO:0004295; F:trypsin activity; IEA.			
DR	GO; GO:0006508; P:proteolysis and peptidolysis; IEA.			
DR	InterPro; IPR000152; Asx_hydroxyl_s			
DR	InterPro; IPR009003; Cys_Ser_trypsin			
DR	InterPro; IPR001881; EGF_Ca			
DR	InterPro; IPR006209; EGF_like			

DR InterPro; IPR002383; GLA blood.
DR InterPro; IPR006210; IEGF.
DR InterPro; IPR001254; Peptidase S1.
DR InterPro; IPR001314; Peptidase_S1A.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00008; EGF; 2.
DR Pfam; PF00594; Gla; 1.
DR Pfam; PF00089; Chymotrypsin.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR SMART; SMC00179; EGF_CA; 1.
DR SMART; SMC00069; GLA; 1.
DR SMART; SMC0020; Tryp_SPC; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS01134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease; Signal.
FT SIGNAL 1 42 POTENTIAL.
FT CHAIN 43 192 PROTEIN C LIGHT CHAIN.
FT CHAIN 193 194 PROTEIN C CONNECTING DIPEPTIDE.
FT CHAIN 195 456 PROTEIN C HEAVY CHAIN.
SQ SEQUENCE 456 AA; 50813 MW; 7AD3A8C1C34E59FF CRC64;

Query Match 72.7%; Score 144; DB 6; Length 456;

Best Local Similarity 61.4%; Pred. No. 9.3e-16;

Matches 27; Conservative 6; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXLLRHSSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44

DB 43 ANSFLLEIRAGSLRECEMEEICDFEAKEIFQNVDDTLAYWSKY 86

RESULT 2

ID Q91WN8 PRELIMINARY; PRT; 460 AA.
AC Q91WN8;
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Similar to protein C.
GN PROC.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Strausberg R.;
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL; BC013896; AAH13896.1; -.
DR HSSP; P00761; 1ANI.
DR MGD; MGI:97771; Proc.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0008235; F:peptidase activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR00152; ASX hydroxyl S.
DR InterPro; IPR009003; Cys_Ser_trypsin.
DR InterPro; IPR01881; EGF_Ca.
DR InterPro; IPR006209; EGF-like.
DR InterPro; IPR002383; GLA_blood.
DR InterPro; IPR01254; Peptidase S1.
DR InterPro; IPR001314; Peptidase_S1A.
DR InterPro; IPR000294; VitK_dep_GLA.

DR Pfam; PF00008; EGF; 2.
DR Pfam; PF00594; Gla; 1.
DR Pfam; PF00089; Chymotrypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SMC00179; EGF_CA; 1.
DR SMART; SMC00069; GLA; 1.
DR SMART; SMC0020; Tryp_SPC; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS01134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
DR EGF-like domain; Hydrolase; Protease; Serine protease.
SQ SEQUENCE 460 AA; 51818 MW; 0117F2E5E8FCC274 CRC64;

Query Match 67.2%; Score 133; DB 11; Length 460;

Best Local Similarity 56.8%; Pred. No. 6.9e-14;

Matches 25; Conservative 7; Mismatches 12; Indels 0; Gaps 0;

QY 1 ANSFLXLLRHSSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44

DB 42 ANSFLLEMRPGSLRECEMEEICDFEAKEIFQNVDDTLAFWKY 85

RESULT 3

Q8J002 PRELIMINARY; PRT; 55 AA.
AC Q8J002;
DT 01-MAR-2003 (TREMBLrel. 23, Created)
DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Protein C (Fragment).
GN PROC.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kinoshita S., Iida H., Inoue S., Watanabe K., Kurihara M., Wada Y.,
RA Ono M., Dongchon K., Hamasaki N.;
RT "Gene Analysis of Anticoagulation Factors in Japanese Thrombotic
RT Patients. Genetic Background of Thrombophilia in Japan."
RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB083700; BAC21172.1; -.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR InterPro; IPR002383; GLA blood.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00594; Gla; 1.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SMC00069; GLA; 1.
FT NON TER 1 55
FT NON TER 55 55
SQ SEQUENCE 55 AA; 6527 MW; 4F89496534A78836 CRC64;

Query Match 65.7%; Score 130; DB 4; Length 55;

Best Local Similarity 70.3%; Pred. No. 2.2e-14;

Matches 26; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXLLRHSSILRXICIXXICDFXXAKXIFEDVDDT 37

DB 19 ANSFLKELRHSSILRECEMEEICDFEAKEIFQNVDDT 55

RESULT 4

Q8IXB5 PRELIMINARY; PRT; 55 AA.
ID Q8IXB5
AC Q8IXB5;

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DT 01-MAR-2003 (T-EMBLrel. 23, Created)
DT 01-MAR-2003 (T-EMBLrel. 23, Last sequence update)
DE 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
DE Protein C (Fragment).
GN PROCI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]_TaxID=9606;
RP SEQUENCE FROM N.A.
RA Hamaaki S., Kang D., Kinoshita S., Iida K., Inoue S., Watanabe K.,
RA Kurihara M., Wada Y., Ono M.;
RT "Gene analysis of anticoagulation factors in Japanese thrombotic
RT patients.Genetic background of thrombophilia in Japan.";
RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB086851; BC53631.1;
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR InterPro; IPR002393; GLA blood.
DR Pfam; PF00594; gla; 1.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00069; GLA; 1.
FT NON_TER 1 55
FT NON_TER 55
SQ SEQUENCE 55 AA; 6475 MW; 3803696534BC9289 CRC64;

Query Match 65.7%; Score 130; DB 4; Length 55;
Best Local Similarity 70.3%; Pred. No. 2,2e-14;
Matches 26; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSLRXICIXICDPXXAKXIFEDVDDT 37
DB 19 ANSFLFELRHSLRECEIEICDPFEAKEIFQNVDDT 55

RESULT 5
Q99PC6 PRELIMINARY; PRT; 460 AA.
AC Q99PC6;
DT 01-JUN-2001 (T-EMBLrel. 17, Created)
DT 01-JUN-2001 (T-EMBLrel. 17, Last sequence update)
DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
DE Anticoagulant protein C.
GN PROC.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]_TaxID=10090;
RP SEQUENCE FROM N.A.
RA STRAIN=C57BL;
RA Korf I.;
RT "Complete sequence of UC72A01.";
RL Submitted (NOV-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL; AF318182; AAK07918.1; -.
DR HSSP; P04070; 1AUF.
DR MGP; MGI:87771; Proc.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0008233; F:peptidase activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR00152; Asx_hydroxyl_S.
DR InterPro; IPR009003; Cys_ser_trypsin.
DR InterPro; IPR001881; EGF Ca.
DR InterPro; IPR006209; EGF like.
DR InterPro; IPR002383; GLA blood.
DR InterPro; IPR001254; Peptidase S1.
DR InterPro; IPR001314; Peptidase_S1A.
DR InterPro; IPR002394; VitK_dep_GLA.
DR InterPro; IPR00594; gla; 1.
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DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00008; EGF; 2.
DR Pfam; PF00594; gla; 1.
DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR SMART; SM00001; GLABLOOD.
DR SMART; SM00179; EGF_CA; 1.
DR SMART; SM00089; GLA; 1.
DR SMART; SM00020; Tryp_SPC; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
KW EGF-like domain; Hydrolase; Protease; Serine protease.
SQ SEQUENCE 460 AA; 51784 MW; 0293BC25E9D3ED16 CRC64;

Query Match 64.1%; Score 127; DB 11; Length 460;
Best Local Similarity 54.5%; Pred. No. 7,2e-13;
Matches 24; Conservative 7; Mismatches 13; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSLRXICIXICDPXXAKXIFEDVDDT 44
DB 42 ANSFLFELRHSLRECEIEICDLEEAQEIFQNVDTLAFWIKY 85

RESULT 6
Q63207 PRELIMINARY; PRT; 482 AA.
AC Q63207;
DT 01-NOV-1996 (T-EMBLrel. 01, Created)
DT 01-NOV-1996 (T-EMBLrel. 01, Last sequence update)
DT 01-OCT-2003 (T-EMBLrel. 25, Last annotation update)
DE Factor X.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]_TaxID=10116;
RP SEQUENCE FROM N.A.
RC STRAIN=Sprague-Dawley;
RX MEDLINE=96093366; PubMed=8578539;
RA Stanton C., Ross R.P., Hutson S., Wallin R.;
RT "Evidence for competition between vitamin K-dependent clotting factors
RT for intracellular processing by the vitamin K-dependent gamma-
RT carboxylase.";
RL Thromb. Res. 80:63-73(1995).
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL; X79807; CAA56202.1; -.
DR PIR; S49075; EXRT.
DR HSSP; P00742; 1XKA.
DR MEROPS; S01.216; -.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0008233; F:peptidase activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR000152; Asx_hydroxyl_S.
DR InterPro; IPR009003; Cys_ser_trypsin.
DR InterPro; IPR000742; EGF_2.
DR InterPro; IPR001881; EGF Ca.
DR InterPro; IPR001438; EGF II.
DR InterPro; IPR006209; EGF like.
DR InterPro; IPR002383; GLA blood.
DR InterPro; IPR001254; Peptidase S1.
DR InterPro; IPR001314; Peptidase_S1A.
DR InterPro; IPR002394; VitK_dep_GLA.
DR Pfam; PF00008; EGF; 2.
DR Pfam; PF00594; gla; 1.
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DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00010; EGFBL00D.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00179; EGF_CA; 1.
DR SMART; SM00069; GLA_1.
DR SMART; SM00020; TRY2_SPC; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
KW EGF-like domain; Hydrolase; Protease; Serine protease.
SQ SEQUENCE 482 AA; 54265 MW; 0284678E3954A698 CRC64;

Query Match 54.0%; Score 107; DB 11; Length 482;
Best Local Similarity 40.9%; Pred. No. 1.9e-09;
Matches 18; Conservative 9; Mismatches 17; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLRKXCIXICDFXXAKXIFEDVDDTLAFWSKH 44
    ||||| : : : : : : : : : : : : : : : : : : :
Db 41 ANSFFFEIKGNLERECVCEICSFEEAREVFEDNKTETFWNKY 84

RESULT 7
Q7SY86
ID Q7SY86 PRELIMINARY; PRT; 455 AA.
AC Q7SY86;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DE Hypothetical protein.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipoidae; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Whole;
RX MEDLINE=22341132; PubMed=12454917;
RA Klein S.L., Strausberg R.L., Wagner L., Pontius J., Clifton S.W.,
RA Richardson P.;
RT "Genetic and genomic tools for Xenopus research: The NIH Xenopus
RL Dev. Dyn. 225:384-391(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Whole;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McSwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Schmutz E.D., Dickson M.C.,
RA Rodriguez A.C., Guichwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
RA Krzywinski M.I., Skalek U., Smalish D.E., Schnerch A., Schein J.E.,
RA Jones S.J., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RL and mouse cDNA sequences.";
Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
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RN [3]
RP SEQUENCE FROM N.A.
RC TISSUE=Whole;
RA Klein S., Strausberg R.;
RL Submitted (JUL-2003) to the EMBL/GenBank/DBJ databases.
KW Hypothetical protein.
SQ SEQUENCE 455 AA; 51811 MW; 07C027ED2B495330 CRC64;

Query Match 51.5%; Score 102; DB 13; Length 455;
Best Local Similarity 47.7%; Pred. No. 1.3e-08;
Matches 21; Conservative 6; Mismatches 17; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLRKXCIXICDFXXAKXIFEDVDDTLAFWSKH 44
    ||||| : : : : : : : : : : : : : : : : : : :
Db 49 AFNFMEEELKPGSLERECIEKCDFEAEFETKEDTLNFWAKY 92

RESULT 8
Q804X5
ID Q804X5 PRELIMINARY; PRT; 433 AA.
AC Q804X5;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Anticoagulant protein C precursor (EC 3.4.21.69).
GN PROC.
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Gallus.
OX NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RA Davidson C.J., Hirt R.P., Lal K., Snell P., Elgar G.,
RA Tuddenham E.G.D., McVey J.H.;
RT "Comparative sequence analysis and molecular evolution of blood
RL coagulation genes from Gallus gallus and Fugu rubripes.";
Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
EMBL; AF465270; AAC33365.1; .
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0016787; F:hydrolase activity; IEA.
DR GO; GO:0003808; F:protein C (activated) activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR000152; Asx_hydroxyl_S.
DR InterPro; IPR009003; Cys_Ser_trypsin.
DR InterPro; IPR00742; EGF_2.
DR InterPro; IPR001881; EGF_CA.
DR InterPro; IPR006209; EGF_like.
DR InterPro; IPR002383; GLA_blood.
DR InterPro; IPR006210; IEGF.
DR InterPro; IPR001254; Peptidase_sl.
DR InterPro; IPR001314; Peptidase_SIA.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00008; EGF; 1.
DR Pfam; PF00594; Gla; 1.
DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00181; EGF; 2.
DR SMART; SM00179; EGF_CA; 1.
DR SMART; SM00069; GLA; 1.
DR SMART; SM00020; TRY2_SPC; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
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DR PROSITE; PS00135; TRYPSIN_SER; 1.
KW Hydrolase.
SQ SEQUENCE 433 AA; 48689 MW; E09DDE56D7DA2A3 CRC64;

Query Match
Best Local Similarity 50.5%; Score 100; DB 13; Length 433;
Matches 21; Conservative 5; Mismatches 18; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSHSLXRCIXXCDFXXKXKIFEDVDDTLAFWSKH 44
||||| : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 40 ANSFLXLRHSHSLXRCIXXCDFXXKXKIFEDVDDTLAFWSKH 83
||||| : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 9
Q7SXH8 PRELIMINARY; PRT; 524 AA.
AC Q7SXH8;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Body;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M.J., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Raha S.S., Loughellano N.A., Peters G.J., Carninci P., Prange C.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Abramson R.D., Mullany S.J.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
RA Krzywinski M.I., Skalska U., Smallus D.E., Schnerch A., Schein J.E.,
RA Jones S.J., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Body;
RA Strausberg R.;
RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC05596; AAH5596.1; -.
KW Hypothetical protein.
SQ SEQUENCE 524 AA; 59560 MW; 1BAAE08119080325 CRC64;

Query Match
Best Local Similarity 49.5%; Score 98; DB 13; Length 524;
Matches 18; Conservative 8; Mismatches 18; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSHSLXRCIXXCDFXXKXKIFEDVDDTLAFWSKH 44
||||| : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 42 ANTVEELKPGNLERECVEECIDHEAREVERVDKTEIFWAKY 85
||||| : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 10
Q8JHC9 PRELIMINARY; PRT; 443 AA.
ID Q8JHC9;
AC Q8JHC9;
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
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DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Coagulation factor VIII.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RT "Comprehensive analysis of blood coagulation pathways in Teleostei:
RT Evolution of coagulation factor genes and identification of zebrafish
RT factor VIII."
EL Blood Cells Mol. Dis. 0:0-0(2002).
CC -!- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL; AF519546; AAM88342.1; -.
DR EMBL; AF515269; AAN71000.1; -.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004363; F:chymotrypsin activity; IEA.
DR GO; GO:0008333; F:peptidase activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR009003; Cys_Ser_trypsin.
DR InterPro; IPR00742; EGF-2.
DR InterPro; IPR01881; EGF_Ca.
DR InterPro; IPR06205; EGF_Like.
DR InterPro; IPR002383; GLA_Blood.
DR InterPro; IPR006210; IEGF.
DR InterPro; IPR01254; Peptidase_S1.
DR InterPro; IPR001314; Peptidase_S1A.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00008; EGF; 1.
DR Pfam; PF00594; GLA; 1.
DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00181; EGF; 2.
DR SMART; SM00179; EGF_CA; 1.
DR SMART; SM00069; GLA; 1.
DR SMART; SM00020; TRYD_SPC; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
DR PROSITE; PS0240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
KW EGF-like domain; Hydrolase; Protease; Serine protease.
SQ SEQUENCE 443 AA; 48823 MW; 2D2504718AE94FF4 CRC64;

Query Match
Best Local Similarity 48.7%; Score 96.5; DB 13; Length 443;
Best Local Similarity 45.2%; Pred. No. 1.1e-07;
Matches 19; Conservative 7; Mismatches 15; Indels 1; Gaps 1;

QY 1 ANSFLXLRHSHSLXRCIXXCDFXXKXKIFEDVDDTLAFW 41
||||| : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 38 ANSGLFLEMKAGNLERECVEECIDYEAREVEFDDRTKQFW 79
||||| : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 11
Q8N2N6 PRELIMINARY; PRT; 231 AA.
ID Q8N2N6;
AC Q8N2N6;
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein FLJ90093.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
```

DR	GO; GO:0004295; F:trypsin activity; IEA.
DR	GO InterPro; IPRO06508; P:proteasome and peptidolysis; IEA.
DR	GO InterPro; IPR000152; Ase_hydroxyl_S
DR	InterPro; IPR009003; Cys_Ser_trypsin.
DR	InterPro; IPR00742; EGF_2.
DR	InterPro; IPR001881; EGF CA.
DR	InterPro; IPR001438; EGF II.
DR	InterPro; IPR006209; EGF like.
DR	InterPro; IPR002383; GLA_blood.
DR	InterPro; IPR006210; IEGF.
DR	InterPro; IPR001254; Peptidase S1.
DR	InterPro; IPR000294; Vitk_dep_GLA.
DR	Pfam; PF00008; EGF; 2.
DR	Pfam; PF00594; Gla; 1.
DR	PRINTS; PR00010; EGFBLOOD.
DR	PRINTS; PR00001; GLABLOOD.
DR	SMART; SMC0181; EGF; 2.
DR	SMART; SMC0179; EGF CA; 1.
DR	SMART; SMC0069; GLA; 1.
DR	PROSITE; PS00010; ASX_HYDROXYL; 1.
DR	PROSITE; PS00022; EGF_1; 1.
DR	PROSITE; PS01186; EGF_2; 2.
DR	PROSITE; PS01187; EGF CA; 1.
DR	PROSITE; PS00011; GLU_CARBOXYLATION; 1.
DR	PROSITE; PS50240; TRYPSIN_DOM; 1.
DR	PROSITE; PS00134; TRYPSIN_HIS; 1.
SQ	SEQUENCE 340 AA; 39359 MW; EE25D6157720811 CRC64;

Query Match 47.0%; Score 93; DB 11; Length 340;
 Best Local Similarity 36.4%; Pred. No. 3, 1e-07;
 Matches 16; Conservative 9; Mismatches 19; Indels 0; Gaps

Qy	1	ANSFLXXLRHSSILRXICIXICDFXXAKXIFPDVDDTLAFWSKH 44 : : : : : : : : : : : :
Dd	53	ANSPFEFKGNLERQMEIEICSYEVEIRFEIDDEKTEYWTKY 96 : : : : : : : : : : : :

RESULT 13

Q7T3B6	PRELIMINARY;	PRT;	434 AA.
ID	Q7T3B6		
AC	Q7T3B6		
DT	01-OCT-2003 (TrEMBLrel. 25, Created)		
DT	01-OCT-2003 (TrEMBLrel. 25, Last sequence update)		
DE	01-OCT-2003 (TrEMBLrel. 25, Last annotation update)		
DE	Hypothetical protein.		
OS	Brachydanio rerio (Zebrafish) (Danio rerio).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
OC	Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;		
OC	Cyprinidae; Danio.		
OX	NCBI_TaxID=7955;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RC	TTSUB=Kidney.		
EX	MEDLINE=22388257; PubMed=12477932;		
RA	Strausberg K.L., Feingold E.A., Grouse L.H., Derge J.G.,		
RA	Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,		
RA	Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,		
RA	Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,		
RA	Diatchenko L., Marushina K., Farmer A.A., Rubin G.M., Hong L.,		
RA	Scapleton M., Soares M.B., Bonaldo M.P., Casavant T.L., Scheetz T.E.,		
RA	Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C.,		
RA	Rana S., Loquellano N.A., Peters G.J., Abramson R.D., Mullany S.J.,		
RA	Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,		
RA	Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,		
RA	Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,		
RA	Fahy J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,		
RA	Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,		
RA	Blakesley A.C., Touchman J.W., Green E.D., Dickson M.C.,		
RA	Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,		
RA	Krzywinski M.I., Skalska U., Smallos D.E., Schnurch A., Schein J.E.,		
RA	Jones S.J., Marra M.A.;		
RT	"Generation and initial analysis of more than 15,000 full-length human		
RT	and mouse cDNA sequences".		
RT	Genomics 26:27-38, 1995.		

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DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
DR PROSITE; PS00040; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
KW EGF-like domain; Hydrolase; Protease; Serine protease; Signal;
KW Plasmid.
FT SIGNAL.
FT CHAIN
SQ SEQUENCE 481 AA; 53986 MW; CF702DE5EP9D97AE CRC64;

Query/Match 47.0%; Score 93; DB 11; Length 481;
Best Local Similarity 36.4%; Pred.No. 4.5e-07;
Matches 16; Conservative 9; Mismatches 19; Indels 0; Gaps 0;

Qy 1 ANSFLLXLRHSLRXCIXXICIDFXKAKXIFEDVDITLAFWSKH 44
      |||||
      41 ANSFPEEPFKGNLRSECMWECISYEVEIRIFEDDEKTYEWTKY 84

Db

RESULT 15
Q99L32
ID Q99L32 PRELIMINARY; PRT; 481 AA.
AC C99L32
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DT Coagulation factor X.
FE F10.
GN GN
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus;
OC NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RA Strausberg R.;
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
RC -!- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
DR EMBL; BC003877; AAH03877.1; -.
DR HSP; P00742.1XKA.
DR MEROPS; S01.216; -.
DR MGD; MGI:103107; F10.
DR GO; GO:0005576; C:extracellular; IEA.
DR GO; GO:0005509; F:calcium ion binding; IEA.
DR GO; GO:0004263; F:chymotrypsin activity; IEA.
DR GO; GO:0008233; F:peptidase activity; IEA.
DR GO; GO:0004295; F:trypsin activity; IEA.
DR GO; GO:0006508; P:proteolysis and peptidolysis; IEA.
DR InterPro; IPR000152; Asx hydroxyl S.
DR InterPro; IPR009003; Cys Ser trypsin.
DR InterPro; IPR000742; EGF_2.
DR InterPro; IPR001881; EGF_CA.
DR InterPro; IPR001438; EGF_11.
DR InterPro; IPR006209; EGF_Like.
DR InterPro; IPR002383; GLA_blood.
DR InterPro; IPR001254; Peptidase S1.
DR InterPro; IPR001314; Peptidase S1A.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00008; EGF_2.
DR Pfam; PF00594; gla; 1.
DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00010; EGF_blood.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SMC00179; EGF_CA; 1.
DR SMART; SMC0069; GLA; 1.
DR SMART; SMC0020; Tryp_SPC; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.

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OM protein - protein search, using sw model

Run on: March 1, 2004, 10:03:28 ; Search time 16.5 Seconds
(without alignments)
137.669 Million cell updates/sec

Title: SEQ1-32GLU-33ASP

Perfect score: 198

Sequence: 1 ANSFLXLRHSLRXRCIXX.....XXAKXIFEDVDTLAFWSKH 44

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents AA.*

- 1: /cgn2_6/ptodata/2/iaa/5A_COMB.pep.*
- 2: /cgn2_6/ptodata/2/iaa/5B_COMB.pep.*
- 3: /cgn2_6/ptodata/2/iaa/6A_COMB.pep.*
- 4: /cgn2_6/ptodata/2/iaa/6B_COMB.pep.*
- 5: /cgn2_6/ptodata/2/iaa/PTUS_COMB.pep.*
- 6: /cgn2_6/ptodata/2/iaa/backfiles.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	180	90.9	44	3	US-08-955-636-19
2	176	88.9	419	4	US-10-182-263-3
3	176	88.9	419	4	US-10-182-263-4
4	176	88.9	419	4	US-10-182-263-5
5	172	86.9	44	3	US-08-955-636-1
6	172	86.9	44	3	US-08-955-636-20
7	172	86.9	44	3	US-08-955-636-21
8	172	86.9	44	3	US-08-955-636-25
9	172	86.9	45	2	US-08-965-832-2
10	172	86.9	419	1	US-08-295-411-1
11	172	86.9	419	2	US-08-955-471-1
12	172	86.9	419	4	US-08-667-570A-3
13	172	86.9	419	4	US-10-182-263-1
14	172	86.9	419	5	PCT-US92-10242-1
15	172	86.9	460	2	US-08-756-506-2
16	172	86.9	460	2	US-08-756-506-4
17	172	86.9	460	6	5270178-13
18	172	86.9	460	6	5270178-14
19	172	86.9	460	6	5270178-15
20	172	86.9	460	6	5270178-16
21	172	86.9	461	4	US-10-182-263-2
22	172	86.9	461	6	5225537-2
23	172	86.9	461	6	5270178-17
24	172	86.9	461	6	5270178-18
25	172	86.9	461	6	5460953-3
26	171	86.4	44	3	US-08-955-636-22
27	168	84.8	44	3	US-08-955-636-24

28	168	84.8	44	3	US-08-955-636-35
29	168	84.8	419	4	US-10-182-263-6
30	159	80.3	42	2	US-08-745-254A-2
31	159	80.3	461	6	5270178-2
32	155	78.3	41	1	US-08-229-280-5
33	150	75.8	42	4	US-09-383-667-8
34	141	71.2	410	3	US-09-065-872-1
35	141	71.2	410	4	US-09-667-570A-1
36	133	67.2	409	3	US-09-065-872-2
37	133	67.2	409	4	US-09-667-570A-2
38	119	60.1	44	3	US-08-955-636-23
39	109	55.1	44	3	US-08-955-636-2
40	106	53.5	139	1	US-08-330-978-2
41	106	53.5	139	1	US-08-474-042-2
42	106	53.5	139	1	US-08-484-558-2
43	106	53.5	139	1	US-08-774-592-2
44	106	53.5	437	1	US-08-487-037-2
45	106	53.5	437	1	US-08-487-037-3

ALIGNMENTS

RESULT 1

US-08-955-636-19
; Sequence 19, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelsestuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 19
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: MOD RES
; LOCATION: (0)..(0)
; OTHER INFORMATION: Xaa-gamma carboxyglutamic acid or glutamic acid
US-08-955-636-19

Query Match 90.9%; Score 180; DB 3; Length 44;
Best Local Similarity 100.0%; Pred. No. 1.4e-23;
Matches 44; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	1	ANSFLXLRHSLRXRCIXXCIFDXKXKXIFEDVDTLAFWSKH 44
DB	1	ANSFLXLRHSLRXRCIXXCIFDXKXKXIFEDVDTLAFWSKH 44

RESULT 2

US-10-182-263-3
; Sequence 3, Application US/10182263
; Patent No. 6630138
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; PRIOR FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1


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; SEQ ID NO 3
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-3

Query Match      88.9%; Score 176; DB 4; Length 419;
Best Local Similarity 77.3%; Pred. No. 9.3e-22;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHSSLRXRCIXXICDFXXAKXIFEDVDDTLAFWSKH 44
Db 1 ANSFLRLRHGSLRECEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 3
US-10-182-263-4
; Sequence 4, Application US/10182263
; Patent No. 6630138
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-4

Query Match      88.9%; Score 176; DB 4; Length 419;
Best Local Similarity 77.3%; Pred. No. 9.3e-22;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHSSLRXRCIXXICDFXXAKXIFEDVDDTLAFWSKH 44
Db 1 ANSFLRLRHGSLRECEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 4
US-10-182-263-5
; Sequence 5, Application US/10182263
; Patent No. 6630138
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 5
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-5

Query Match      88.9%; Score 176; DB 4; Length 419;
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Best Local Similarity 77.3%; Pred. No. 9.3e-22;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHSSLRXRCIXXICDFXXAKXIFEDVDDTLAFWSKH 44
Db 1 ANSFLRLRHGSLRECEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 5
US-08-955-636-1
; Sequence 1, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelsestuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; FILE REFERENCE: POLYPEPTIDES
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 1
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; NAME/KEY: MOD_RES
; LOCATION: (0)...(0)
; OTHER INFORMATION: Xaa-gamma carboxyglutamic acid or glutamic acid
US-08-955-636-1

Query Match      86.9%; Score 172; DB 3; Length 44;
Best Local Similarity 95.5%; Pred. No. 3.3e-22;
Matches 42; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHSSLRXRCIXXICDFXXAKXIFEDVDDTLAFWSKH 44
Db 1 ANSFLXLRHSSLRXRCIXXICDFXXAKXIFQNVDDTLAFWSKH 44

RESULT 6
US-08-955-636-20
; Sequence 20, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelsestuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; FILE REFERENCE: POLYPEPTIDES
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSEQ for Windows Version 3.0
; SEQ ID NO 20
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; NAME/KEY: MOD_RES
; LOCATION: (0)...(0)
; OTHER INFORMATION: Xaa-gamma carboxyglutamic acid or glutamic acid
US-08-955-636-20

Query Match      86.9%; Score 172; DB 3; Length 44;
Best Local Similarity 97.7%; Pred. No. 3.3e-22;
Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 1 ANSFLXLRHSSLRXRCIXXICDFXXAKXIFEDVDDTLAFWSKH 44

RESULT 7
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08-955-636-21
Sequence 21, Application US/08955636A
Patent No. 6017882

GENERAL INFORMATION:
APPLICANT: Nelsestuen, Gary
TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
POLYPEPTIDES
FILE REFERENCE: 09531/002001
CURRENT APPLICATION NUMBER: US/08/955,636A
CURRENT FILING DATE: 1997-10-23
NUMBER OF SEQ ID NOS: 35
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 21
LENGTH: 44
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: MOD RES
LOCATION: (0)...(0)
OTHER INFORMATION: Xaa-gamma carboxyglutamic acid or glutamic acid
-08-955-636-21

Query Match 86.9%; Score 172; DB 3; Length 44;
Best Local Similarity 97.7%; Pred. No. 3.3e-22;
Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

1 ANSFLXLLRHSLRXKXICIXXICDFXXAKXIFEDVDDTLAPWSKH 44
|||||
1 ANSFLXLLRHSLRXKXICIXXICDFXXAKXIFEDVDDTLAPWSKH 44

RESULT 8

US-08-955-636-25
Sequence 25, Application US/08955636A
Patent No. 6017882
GENERAL INFORMATION:

APPLICANT: Nelsestuen, Gary
TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
POLYPEPTIDES
FILE REFERENCE: 09531/002001
CURRENT APPLICATION NUMBER: US/08/955,636A
CURRENT FILING DATE: 1997-10-23
NUMBER OF SEQ ID NOS: 35
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 25
LENGTH: 44
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: MOD RES
LOCATION: (0)...(0)
OTHER INFORMATION: Xaa-gamma carboxyglutamic acid or glutamic acid
US-08-955-636-25

Query Match 86.9%; Score 172; DB 3; Length 44;
Best Local Similarity 97.7%; Pred. No. 3.3e-22;
Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ANSFLXLLRHSLRXKXICIXXICDFXXAKXIFEDVDDTLAPWSKH 44
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Db 1 ANSFLXLLRHSLRXKXICIXXICDFXXAKXIFEDVDDTLAPWSKH 44

RESULT 9

US-08-965-832-2
Sequence 2, Application US/08965832
Patent No. 5847095
GENERAL INFORMATION:
APPLICANT: CHARLES T. ESMON AND MIKHAIL D. SMIRNOV
TITLE OF INVENTION: Modified Protein C
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Patrea L. Pabst

STREET: 2800 One Atlantic Center, 1201 West
STREET: Peachtree Street
CITY: Atlanta
STATE: GA
COUNTRY: USA
ZIP: 30309-3450
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION NUMBER: US/08/965,832
FILING DATE: 7-NOV-1997
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/745,254
FILING DATE: 8-NOV-1996
PRIOR APPLICATION NUMBER:
APPLICATION NUMBER: 60/053,768
FILING DATE: 25-JUL-1997
ATTORNEY/AGENT INFORMATION:
NAME: Pabst, Patrea L.
REGISTRATION NUMBER: 31,284
REFERENCE/DOCKET NUMBER: OMRF 165/167
TELECOMMUNICATION INFORMATION:
TELEPHONE: (404)-873-8794
TELEFAX: (404)-873-8795
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 45 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY:
LOCATION: 6, 7, 14, 16, 19, 20, 25, 26, 29
OTHER INFORMATION: /note= "where Xaa means gamma
OTHER INFORMATION: carboxylglutamic acid"
FEATURE:
NAME/KEY:
LOCATION:
OTHER INFORMATION: /note= "partial sequence of human protein C"
US-08-965-832-2

Query Match 86.9%; Score 172; DB 2; Length 45;
Best Local Similarity 95.5%; Pred. No. 3.3e-22;
Matches 42; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 ANSFLXLLRHSLRXKXICIXXICDFXXAKXIFEDVDDTLAPWSKH 44
|||||
Db 1 ANSFLXLLRHSLRXKXICIXXICDFXXAKXIFEDVDDTLAPWSKH 44

RESULT 10

US-08-295-411-1
Sequence 1, Application US/08295411
Patent No. 5679639
GENERAL INFORMATION:
APPLICANT: Griffin, John H.
APPLICANT: Masters, Rolf M.
TITLE OF INVENTION: Serine Protease-Derived Polypeptides and
TITLE OF INVENTION: Anti-Peptide Antibodies, Systems and Therapeutic Methods
TITLE OF INVENTION: for Inhibiting Coagulation
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Office of Patent Counsel, The Scripps
ADDRESSEE: Research Institute
STREET: 10666 No. 5679639th Torrey Pines Road, TPC 8
CITY: La Jolla
STATE: CA
COUNTRY: USA

```

ZIP: 92037
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION NUMBER: US/08/955,471
FILING DATE: 22-AUG-1994
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/793,989
FILING DATE: 18-NOV-1991
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: Fitting, Thomas
REGISTRATION NUMBER: 34,163
REFERENCE/DOCKET NUMBER: TSRI263.0C1
TELEPHONE: 619-554-2937
TELEFAX: 619-554-6312
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 419 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: Region
LOCATION: 1..157
OTHER INFORMATION: /note= "Protein C Light Chain"
FEATURE:
NAME/KEY: Region
LOCATION: 158..169
OTHER INFORMATION: /note= "Protein C Activation"
OTHER INFORMATION: Peptide"
FEATURE:
NAME/KEY: Region
LOCATION: 170..419
OTHER INFORMATION: /note= "Protein C Heavy Chain"
US-08-295-411-1
Query Match 86.9%; Score 172; DB 1; Length 419;
Best Local Similarity 75.0%; Pred. No. 4,5e-21;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;
QY 1 ANSFLXLRHSSLRXCIXICDFXAXKIFEDVDDTLAFWSKH 44
DB 1 ANSFLRLHSSLRERCIEICDFEAKEIFQNVDDTLAFWSKH 44
RESULT 11
US-08-955-471-1
Sequence 1, Application US/08955471
Patent No. 5968751
GENERAL INFORMATION:
APPLICANT: Griffin, John H.
APPLICANT: Masters, Rolf M.
TITLE OF INVENTION: Serine Protease-Derived Polypeptides and
TITLE OF INVENTION: Anti-Peptide Antibodies, Systems and Therapeutic Methods
TITLE OF INVENTION: for Inhibiting Coagulation
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Office of Patent Counsel, The Scripps
ADDRESSEE: Research Institute
STREET: 10666 No. 5968/5th Torrey Pines Road, TPC 8
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION NUMBER: US/08/955,471
FILING DATE: 22-AUG-1994
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/793,989
FILING DATE: 18-NOV-1991
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: Fitting, Thomas
REGISTRATION NUMBER: 34,163
REFERENCE/DOCKET NUMBER: TSRI263.0C1
TELEPHONE: 619-554-2937
TELEFAX: 619-554-6312
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 419 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: Region
LOCATION: 1..157
OTHER INFORMATION: /note= "Protein C Light Chain"
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LOCATION: 158..169
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OTHER INFORMATION: Peptide"
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LOCATION: 170..419
OTHER INFORMATION: /note= "Protein C Heavy Chain"
US-08-295-411-1
Query Match 86.9%; Score 172; DB 1; Length 419;
Best Local Similarity 75.0%; Pred. No. 4,5e-21;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;
QY 1 ANSFLXLRHSSLRXCIXICDFXAXKIFEDVDDTLAFWSKH 44
DB 1 ANSFLRLHSSLRERCIEICDFEAKEIFQNVDDTLAFWSKH 44
RESULT 12
US-09-667-570A-3
Sequence 3, Application US/09667570A
Patent No. 6436397
GENERAL INFORMATION:
APPLICANT: Baker, Jeffrey C
APPLICANT: Carlson, Andrew D
APPLICANT: Huang, Litua
APPLICANT: Sheliga, Theodore A
TITLE OF INVENTION: Improved Methods for Processing Activated Protein C
FILE REFERENCE: X-11796A
CURRENT APPLICATION NUMBER: US/09/667,570A
CURRENT FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: 60/045,255
PRIOR FILING DATE: 1997-04-28
NUMBER OF SEQ ID NOS: 3
SOFTWARE: Patent in version 3.1
SEQ ID NO 3
LENGTH: 419
TYPE: PRT
ORGANISM: Homo sapiens
US-09-667-570A-3

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100

TELECOMMUNICATION INFORMATION:
 TELEPHONE: 206-442-6672
 TELEFAX: 206-442-6678
 INFORMATION FOR SEQ ID NO: 2:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 460 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-756-506-2

Query Match 86.9%; Score 172; DB 2; Length 460;
 Best Local Similarity 75.0%; Fred. No. 5e-21;
 Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLRXCIXICDFXXAKXIFEDVDDTLAPWSKH 44
 Db 43 ANSFLXLRHSSLRXCIXICDFXXAKXIFEDVDDTLAPWSKH 86

Search completed: March 1, 2004, 10:12:20
 Job time: 17.5 secs

(Handwritten signatures and scribbles)

GenCore version 5.1.6
Copyright (C) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: March 1, 2004, 10:01:28 ; Search time 28 Seconds
(without alignments)
331.812 Million cell updates/sec

Title: SEQ1-32GLU-33ASP
Perfect score: 198
Sequence: 1 ANSFLXLRHSLRXRCIXX.....XXAKXIFEDVDDTLAFWSKH 44

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 809742 seqs, 211153259 residues

Total number of hits satisfying chosen parameters: 809742

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA:

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- 2: /cgn2_6/ptodata/2/pubpaa/PCT_NEW_PUB.pep.*
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- 18: /cgn2_6/ptodata/2/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	176	88.9	419	14	US-10-182-263-3
2	176	88.9	419	14	US-10-182-263-4
3	176	88.9	419	14	US-10-182-263-5
4	176	88.9	419	15	US-10-168-407-3
5	176	88.9	419	15	US-10-168-407-4
6	172	86.9	44	14	US-10-298-330-1
7	172	86.9	419	10	US-09-978-917A-4
8	172	86.9	419	14	US-10-182-263-1
9	172	86.9	419	15	US-10-168-407-1
10	172	86.9	461	10	US-09-978-917A-2
11	172	86.9	461	14	US-10-182-263-2
12	172	86.9	461	15	US-10-168-407-2
13	168	84.8	419	14	US-10-182-263-6
14	168	84.8	419	15	US-10-168-407-5
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16	109	55.1	44	14	US-10-298-330-2	Sequence 2, Appli
17	102	51.5	139	15	US-10-360-101-232	Sequence 232, App
18	102	51.5	488	14	US-10-348-504-44	Sequence 44, Appl
19	102	51.5	488	14	US-10-407-123-27	Sequence 27, Appl
20	95	48.0	44	14	US-10-298-330-18	Sequence 18, Appl
21	92	46.5	406	10	US-09-782-587B-3	Sequence 3, Appli
22	92	46.5	406	15	US-10-383-898-1	Sequence 1, Appli
23	92	46.5	466	14	US-10-017-122-2	Sequence 2, Appli
24	92	46.5	466	15	US-10-375-741-14	Sequence 14, Appl
25	91	46.0	42	16	US-10-038-854-97	Sequence 97, Appl
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32	88	44.4	405	15	US-10-360-101-225	Sequence 225, App
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34	80	40.4	40	14	US-10-298-330-23	Sequence 23, Appl
35	78	39.4	40	14	US-10-298-330-22	Sequence 22, Appl
36	78	39.4	40	14	US-10-298-330-25	Sequence 25, Appl
37	78	39.4	96	10	US-09-759-130B-313	Sequence 313, App
38	78	39.4	96	14	US-10-189-123-43	Sequence 43, Appl
39	78	39.4	96	14	US-10-188-495-43	Sequence 43, Appl
40	78	39.4	209	10	US-09-759-130B-312	Sequence 312, App
41	78	39.4	209	14	US-10-189-123-42	Sequence 42, Appl
42	78	39.4	209	14	US-10-188-495-42	Sequence 42, Appl
43	78	39.4	226	10	US-09-759-130B-310	Sequence 310, App
44	78	39.4	226	14	US-10-189-123-40	Sequence 40, Appl
45	78	39.4	226	14	US-10-188-495-40	Sequence 40, Appl

ALIGNMENTS

RESULT 1
US-10-182-263-3
; Sequence 3, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Garlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 3
; LENGTH: 419
; TYPE: PPT
; ORGANISM: Homo sapiens
US-10-182-263-3

Query Match 88.9%; Score 176; DB 14; Length 419;
Best Local Similarity 77.3%; Pred. No. 5,6e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
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DB 1 ANSFLXLRHSLRXRCIXXICDFXXAKXIFEDVDDTLAFWSKH 44

RESULT 2
US-10-182-263-4
; Sequence 4, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:

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; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-4

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Best Local Similarity 77.3%; Pred. No. 5.6e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

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US-10-182-263-5
; Sequence 5, Application US/10182263
; Publication No. US2003002354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 5
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-5

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Best Local Similarity 77.3%; Pred. No. 5.6e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

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RESULT 4
US-10-168-407-3
; Sequence 3, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
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; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 3
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-3

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Best Local Similarity 77.3%; Pred. No. 5.6e-21;
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US-10-168-407-4
; Sequence 4, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-4

Query Match      88.9%; Score 176; DB 15; Length 419;
Best Local Similarity 77.3%; Pred. No. 5.6e-21;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

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RESULT 6
US-10-298-330-1
; Sequence 1, Application US/10298330
; Publication No. US20030100506A1
; GENERAL INFORMATION:
; APPLICANT: Nelsestuen, Gary L.
; TITLE OF INVENTION: Modified Vitamin K-Dependent
; FILE REFERENCE: 09531-127001
; CURRENT APPLICATION NUMBER: US/10/298,330
; CURRENT FILING DATE: 2002-11-18
; PRIOR APPLICATION NUMBER: 09/497,591
; PRIOR FILING DATE: 2000-02-03
; PRIOR APPLICATION NUMBER: 09/302,239
; PRIOR FILING DATE: 1999-04-29
; PRIOR APPLICATION NUMBER: 08/955,636
; PRIOR FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 27
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: VARIANT
; LOCATION: 6, 7, 14, 16, 19, 20, 25, 26, 29
; OTHER INFORMATION: Xaa = gamma carboxyglutamic or glutamic acid
US-10-298-330-1
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Best Local Similarity 95.5%; Pred. No. 2.2e-21;
Matches 42; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

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RESULT 7
US-09-978-917A-4
; Sequence 4, Application US/09978917A
; Publication No. US20030027299A1
; GENERAL INFORMATION:
; APPLICANT: Maxygen Aps; Maxygen Holdings
; TITLE OF INVENTION: Protein C or activated protein C-like molecules
; FILE REFERENCE: 0219us310 - protein C
; CURRENT APPLICATION NUMBER: US/09/978,917A
; CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-978-917A-4

Query Match      86.9%; Score 172; DB 10; Length 419;
Best Local Similarity 75.0%; Pred. No. 2.6e-20;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

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RESULT 8
US-10-182-263-1
; Sequence 1, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 1
; LENGTH: 419
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; ORGANISM: Homo sapiens
US-10-182-263-1

Query Match      86.9%; Score 172; DB 14; Length 419;
Best Local Similarity 75.0%; Pred. No. 2.6e-20;
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RESULT 9
US-10-168-407-1
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; Publication No. US20030207435A1
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; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: Patentin version 3.1
; SEQ ID NO 1
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-1

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Best Local Similarity 75.0%; Pred. No. 2.6e-20;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

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DB 1 ANSFLRHSLSRECEIEICDFEAKEIFQNVDDTLAFWSKH 44
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RESULT 10
US-09-978-917A-2
; Sequence 2, Application US/09978917A
; Publication No. US20030027299A1
; GENERAL INFORMATION:
; APPLICANT: Maxygen Aps; Maxygen Holdings
; TITLE OF INVENTION: Protein C or activated protein C-like molecules
; FILE REFERENCE: 0219us310 - protein C
; CURRENT APPLICATION NUMBER: US/09/978,917A
; CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: Patentin Ver. 2.1
; SEQ ID NO 2
; LENGTH: 461
; TYPE: PRT
; ORGANISM: Homo sapiens
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; LOCATION: (1)....(42)
; FEATURE:
; NAME/KEY: CHAIN
; LOCATION: (43)....(461)
US-09-978-917A-2

Query Match      86.9%; Score 172; DB 10; Length 461;
Best Local Similarity 75.0%; Pred. No. 2.9e-20;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

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DB 43 ANSFLRHSLSRECEIEICDFEAKEIFQNVDDTLAFWSKH 86
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RESULT 11
US-10-182-263-2
; Sequence 2, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
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; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 461
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; ORGANISM: Homo sapiens
US-10-182-263-2

Query Match 86.9%; Score 172; DB 14; Length 461;
Best Local Similarity 75.0%; Pred. No. 2.9e-20;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

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Db 43 ANSFLELRQGSLEERCIEICDFFEAKEIFQVDDTLAFWSKH 86
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RESULT 12

US-10-168-407-2
; Sequence 2, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:

; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2
; LENGTH: 461
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-2

Query Match 86.9%; Score 172; DB 15; Length 461;
Best Local Similarity 75.0%; Pred. No. 2.9e-20;
Matches 33; Conservative 2; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLRXCIXXICDFFXKXKIFEDVDDTLAFWSKH 44
|||||
Db 43 ANSFLELRHSSLEERCIEICDFFEAKEIFQVDDTLAFWSKH 86
|||||

RESULT 13

US-10-182-263-6
; Sequence 6, Application US/10182263
; Publication No. US2003022354A1
; GENERAL INFORMATION:

; APPLICANT: Jones, Bruce E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-07-22
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 6
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-6

Query Match 84.8%; Score 168; DB 14; Length 419;
Best Local Similarity 75.0%; Pred. No. 1.2e-19;
Matches 33; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLRXCIXXICDFFXKXKIFEDVDDTLAFWSKH 44
|||||
Db 1 ANSFLELRQGSLEERCIEICDFFEAKEIFEDVDDTLAFWSKH 44
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RESULT 14

US-10-168-407-5
; Sequence 5, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:

; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 5
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-5

Query Match 84.8%; Score 168; DB 15; Length 419;
Best Local Similarity 75.0%; Pred. No. 1.2e-19;
Matches 33; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLRXCIXXICDFFXKXKIFEDVDDTLAFWSKH 44
|||||
Db 1 ANSFLELRQGSLEERCIEICDFFEAKEIFEDVDDTLAFWSKH 44
|||||

RESULT 15

US-10-168-407-6
; Sequence 6, Application US/10168407
; Publication No. US20030207435A1
; GENERAL INFORMATION:

; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13610
; CURRENT APPLICATION NUMBER: US/10/168,407
; CURRENT FILING DATE: 2002-11-04
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 6
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-168-407-6

Query Match 84.8%; Score 168; DB 15; Length 419;
Best Local Similarity 75.0%; Pred. No. 1.2e-19;
Matches 33; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXLRHSSLRXCIXXICDFFXKXKIFEDVDDTLAFWSKH 44
|||||
Db 1 ANSFLELRQGSLEERCIEICDFFEAKEIFEDVDDTLAFWSKH 44
|||||

Search completed: March 1, 2004, 10:11:10
Job time : 30 secs

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OM protein - protein search, using sw model

Run on: March 1, 2004, 09:54:37 ; Search time 50 Seconds
(without alignments)
248.642 Million cell updates/sec

Title: SEQ1-4SUBS

Perfect score: 197

Sequence: 1 ANGFLXLRGSLKRCXCIXX.....XXAKXIFedVDDTLAFWSKH 44

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_29Jan04:*
1: Geneseq1980s:*
2: Geneseq1990s:*
3: Geneseq2000s:*
4: Geneseq2001s:*
5: Geneseq2002s:*
6: Geneseq2003as:*
7: Geneseq2003bs:*
8: Geneseq2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	179	90.9	44	2 AAY18300	Modified
2	179	90.9	419	4 AAB82677	Human pro
3	179	90.9	419	4 AAB82678	Human pro
4	179	90.9	419	4 AAE08630	Human pro
5	176	89.3	44	2 AAY18301	Modified
6	174	88.3	419	4 AAB82676	Human pro
7	174	88.3	419	4 AAB82675	Human pro
8	174	88.3	419	4 AAE08628	Human pro
9	174	88.3	419	4 AAE08627	Human pro
10	174	88.3	419	4 AAE08629	Human pro
11	173	87.8	44	2 AAY18298	Modified
12	170	86.3	44	2 AAY18299	Modified
13	168	85.3	44	2 AAY18297	Modified
14	168	85.3	44	2 AAY18307	Modified
15	168	85.3	45	5 ABB79949	Human pro
16	164	83.2	45	5 ABB79946	Human pro
17	164	83.2	45	7 ADB71159	Human pro
18	162	82.2	45	5 ABB79950	Human pro
19	160	81.2	44	2 AAY18309	Modified
20	160	81.2	44	2 AAY18303	Human pro
21	160	81.2	44	4 AAB36402	Human pro
22	160	81.2	44	7 ADD50094	Human vit
23	160	81.2	45	2 AAW5710	Partial h
24	160	81.2	45	5 ABB79947	Human pro
25	160	81.2	45	7 ADB71155	Human pro

26	160	81.2	415	3 AAY56803	Truncated
27	160	81.2	419	2 AAR35760	Protein C
28	160	81.2	419	2 AAW72753	Primary s
29	160	81.2	419	4 AAB82673	Wild-type
30	160	81.2	419	4 AAB36896	Human pro
31	160	81.2	419	4 AAB36897	Human pro
32	160	81.2	419	4 AAB36898	Human pro
33	160	81.2	419	4 AAB36894	Human pro
34	160	81.2	419	4 AAE08625	Human mat
35	160	81.2	419	5 AAU99005	Human pro
36	160	81.2	419	5 AAU99006	Human pro
37	160	81.2	419	5 AAU99008	Human pro
38	160	81.2	419	5 AAU99018	Human pro
39	160	81.2	419	5 AAU99026	Human pro
40	160	81.2	419	5 AAU99037	Human pro
41	160	81.2	419	5 AAU99049	Human pro
42	160	81.2	419	5 AAU99063	Human pro
43	160	81.2	419	5 AAU99072	Human pro
44	160	81.2	419	5 AAU99083	Human pro
45	160	81.2	419	5 AAU99084	Human pro

ALIGNMENTS

RESULT 1
ID AAY18300 standard; peptide; 44 AA.
XX
AC AAY18300;
XX
DT 17-AUG-1999 (first entry)
XX
DE Modified GLA domain of vitamin K-dependent protein.
XX
KW GLA domain; muten; vitamin K-dependent protein; clotting disorder;
KW therapy.
XX
OS Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Misc-difference 1...44 /note= "Xaa= gamma-carboxyglutamic acid, or glutamic acid"
FT
XX
PN WO9920767-A1.
XX
PD 29-APR-1999.
XX
PF 20-OCT-1998; 98WO-US022152.
XX
PR 23-OCT-1997; 97US-00955636.
XX
PA (MINU) UNIV MINNESOTA.
XX
PI Nelsestuen GL;
XX
DR WPI; 1999-288309/24.
XX
PT Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid domain, useful for treating clotting disorders.
XX
PS Claim 9; Page 79; 86pp; English.
XX
CC This sequence represents a modified GLA (gamma-carboxyglutamic acid) domain. The invention relates to a vitamin K-dependent polypeptide comprising a modified GLA domain containing an amino acid substitution which enhances membrane binding of the modified polypeptide as compared to the native polypeptide. The polypeptide is used to treat a clotting disorder by decreasing or increasing clot formation. Modification of the GLA domain results in a protein which has enhanced membrane binding affinity as compared to the native protein

```
XX SQ Sequence 44 AA;
Query Match 90.9%; Score 179; DB 2; Length 44;
Best Local Similarity 100.0%; Pred. No. 2.3e-22;
Matches 44; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ANSFLXXLRQGSIXRXCIXXICDPFXAKXIFEDVDDTLAFWSKH 44
DB 1 ANSFLXXLRQGSIXRXCIXXICDPFXAKXIFEDVDDTLAFWSKH 44

RESULT 2
AAB82677
ID AAB82677 standard; protein; 419 AA.
XX
AC AAB82677;
XX
DT 15-OCT-2001 (first entry)
XX
DE Human protein C derivative (H10Q/S11G/Q32E/N33D/L194S).
XX
KW Protein C; human; coronary syndrome; thrombosis; angina;
KW myocardial infarction; vascular occlusive disorder; hypercoagulation;
KW sepsis; protein C deficiency; occlusion; thromboembolism; stenosis;
KW antibacterial; immunosuppressive; thrombolytic; cardiac; antiangiinal;
KW anticoagulant; therapy; mutant; muten.
XX
OS Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Domain 1..45 /note= "Gla domain"
FT Modified-site 6 /note= "gamma-carboxylated"
FT Modified-site 7 /note= "gamma-carboxylated"
FT Misc-difference 10 /note= "gamma-carboxylated"
FT Misc-difference 11 /note= "His in wild-type protein"
FT Modified-site 14 /note= "Ser in wild-type protein"
FT Modified-site 16 /note= "gamma-carboxylated"
FT Modified-site 19 /note= "gamma-carboxylated"
FT Modified-site 20 /note= "gamma-carboxylated"
FT Modified-site 25 /note= "gamma-carboxylated"
FT Modified-site 26 /note= "gamma-carboxylated"
FT Modified-site 29 /note= "gamma-carboxylated"
FT Misc-difference 32 /note= "N-glycosylated"
FT Misc-difference 33 /note= "Gln in wild-type protein"
FT Disulfide-bond 50..59 /note= "Asn in wild-type protein"
FT Disulfide-bond 59..64
FT Disulfide-bond 80..89
FT Disulfide-bond 98..109
FT Disulfide-bond 120..133
FT Disulfide-bond 141..277
FT Cleavage-site 156..157
/note= "cleavage makes a 2-chain inactive precursor (155-
amino acid light chain attached via a disulfide bond to a
262-amino acid heavy chain)"
FT Peptide 158..169
FT /note= "activation peptide; removal activates the 2-chain
zymogen"
FT
```

```
FT Cleavage-site 169..170
/note= "thrombin cleavage site"
FT Misc-difference 194
/note= "Leu in wild-type protein"
FT Disulfide-bond 196..212
FT Modified-site 248 /note= "N-glycosylated"
FT Modified-site 313 /note= "N-glycosylated"
FT Modified-site 329 /note= "N-glycosylated"
FT Disulfide-bond 331..345
FT Disulfide-bond 356..384
XX WO200157193-A2.
XX
XX 09-AUG-2001.
XX
XX 19-JAN-2001; 2001WO-US000020.
XX
XX 02-FEB-2000; 2000US-0179801P.
XX 14-MAR-2000; 2000US-0189197P.
XX (ELIL ) LILLY & CO ELI.
XX Gerlitz BE, Jones BE;
XX WPI; 2001-496919/54.
XX N-PSDB; AAB26365.
XX
XX Novel human protein C derivative for treating, e.g., myocardial
infarction, unstable angina, sepsis, thrombotic disorders, acute arterial
thrombotic occlusion, and thromboembolism.
XX
XX Claim 5; Page 54-55; 63pp; English.
XX
XX The present sequence is that of a claimed human protein C derivative in
which His at position 10 of the mature wild-type protein C sequence (see
AAB82673) is substituted with Gln, Ser at position 11 with Gly, Gln at
position 32 with Glu, Asn at position 33 with Asp, and Leu at position
194 with Ser. It is an example of protein C derivatives of the invention
that have at least 2 amino acid substitutions, but which have increased
anticoagulant activity and resistance to inactivation by serpins compared
with the wild-type protein, while retaining the biological activity of
the wild-type protein. A method of producing the derivatives using
recombinant DNA methods is claimed. The protein C derivatives are useful
for treating coronary syndromes and disease states predisposing to
thrombosis (e.g. myocardial infarction and unstable angina), vascular
occlusive disorders and hypercoagulable states, sepsis (in combination
with bactericidal permeability increasing protein or with tissue factor
pathway inhibitor), thrombotic disorders (in combination with an anti-
platelet agent or by local delivery through an intracoronary catheter),
protein C deficiency, acute arterial thrombotic occlusion,
thromboembolism, or stenosis in coronary, cerebral or peripheral arteries
or in vascular grafts. Human patients with genetically predisposed
prothrombotic disorders may be treated by gene therapy (all claimed)
XX
XX SQ Sequence 419 AA;
```

```
Query Match 90.9%; Score 179; DB 4; Length 419;
Best Local Similarity 79.5%; Pred. No. 2.5e-21;
Matches 35; Conservative 0; Mismatches 9; Indels 0; Gaps 0;
QY 1 ANSFLXXLRQGSIXRXCIXXICDPFXAKXIFEDVDDTLAFWSKH 44
DB 1 ANSFLXXLRQGSIXRXCIXXICDPFXAKXIFEDVDDTLAFWSKH 44

RESULT 3
AAB82678
ID AAB82678 standard; protein; 419 AA.
XX
XX AAB82678;
AC AAB82678;
```

XX 15-OCT-2001 (first entry)
 XX Human protein C derivative (H10Q/S11G/Q32E/N33D/L194S/T254S).
 DE
 XX
 KW Protein C; human; coronary syndrome; thrombosis; angina;
 KW myocardial infarction; vascular occlusive disorder; hypercoagulation;
 KW sepsis; protein C deficiency; occlusion; thromboembolism; stenosis;
 KW antibacterial; immunosuppressive; thrombolytic; cardiac; antianginal;
 KW anticoagulant; therapy; mutant; mutein.
 XX
 OS Homo sapiens.
 OS Synthetic.
 XX
 XX Key Location/Qualifiers
 XX Domain 1..45
 XX Modified-site /note= "Gla domain"
 XX Modified-site /note= "gamma-carboxylated"
 XX Modified-site /note= "gamma-carboxylated"
 XX Misc-difference 10 /note= "gamma-carboxylated"
 XX Misc-difference 11 /note= "His in wild-type protein"
 XX Modified-site 14 /note= "Ser in wild-type protein"
 XX Modified-site 16 /note= "gamma-carboxylated"
 XX Modified-site 19 /note= "gamma-carboxylated"
 XX Modified-site 20 /note= "gamma-carboxylated"
 XX Modified-site 25 /note= "gamma-carboxylated"
 XX Modified-site 26 /note= "gamma-carboxylated"
 XX Modified-site 29 /note= "gamma-carboxylated"
 XX Misc-difference 32 /note= "N-glycosylated"
 XX Misc-difference 33 /note= "Gln in wild-type protein"
 XX Disulfide-bond 50..69 /note= "Asn in wild-type protein"
 XX Disulfide-bond 59..64
 XX Disulfide-bond 80..89
 XX Disulfide-bond 98..109
 XX Disulfide-bond 120..133
 XX Disulfide-bond 141..277
 XX Cleavage-site 156..157
 XX /note= "cleavage makes a 2-chain inactive precursor (155-
 XX amino acid light chain attached via a disulfide bond to a
 XX 262-amino acid heavy chain)"
 XX Peptide 158..169
 XX /note= "activation peptide; removal activates the 2-chain
 XX zymogen"
 XX Cleavage-site 169..170
 XX Misc-difference 194 /note= "thrombin cleavage site"
 XX Disulfide-bond 196..212 /note= "Leu in wild-type protein"
 XX Modified-site 248
 XX Misc-difference 254 /note= "N-glycosylated"
 XX Modified-site 313 /note= "Thr in wild-type protein"
 XX Modified-site 329 /note= "N-glycosylated"
 XX Disulfide-bond 331..345 /note= "N-glycosylated"
 XX Disulfide-bond 356..384
 XX WO200157193-A2.

XX 09-AUG-2001.
 XX 19-JAN-2001; 2001WO-US000020.
 XX 02-FEB-2000; 2000US-0179801P.
 XX 14-MAR-2000; 2000US-0189197P.
 XX (BLIL) LILLY & CO ELI.
 XX Gerlitz BE, Jones BE;
 XX WPI; 2001-496919/54.
 XX Novel human protein C derivative for treating, e.g., myocardial
 XX infarction, unstable angina, sepsis, thrombotic disorders, acute arterial
 XX thrombotic occlusion, and thromboembolism.
 XX Claim 6; Page 56-57; 63pp; English.
 XX The present sequence is that of a claimed human protein C derivative in
 XX which His at position 10 of the wild-type protein C sequence (see
 XX AAB82673) is substituted with Gln, Ser at position 11 with Gly, Gln at
 XX position 32 with Glu, Asn at position 33 with Asp, Leu at position 194
 XX with Ser, and Thr at position 254 with Ser. It is an example of protein C
 XX derivatives of the invention that have at least 2 amino acid
 XX substitutions, but which have increased anticoagulant activity and
 XX resistance to inactivation by serpins compared with the wild-type
 XX protein, while retaining the biological activity of the wild-type
 XX protein. A method of producing the derivatives using recombinant DNA
 XX methods is claimed. The protein C derivatives are useful for treating
 XX coronary syndromes and disease states predisposing to thrombosis (e.g.
 XX myocardial infarction and unstable angina), vascular occlusive disorders
 XX and hypercoagulable states, sepsis (in combination with bactericidal
 XX permeability increasing protein or with tissue factor pathway inhibitor),
 XX thrombotic disorders (in combination with an anti-platelet agent or by
 XX local delivery through an intracoronary catheter), protein C deficiency,
 XX acute arterial thrombotic occlusion, thromboembolism, or stenosis in
 XX coronary; cerebral or peripheral arteries or in vascular grafts. Human
 XX patients with genetically predisposed prothrombotic disorders may be
 XX treated by gene therapy (all claimed)
 XX Sequence 419 AA;
 XX
 XX Query Match 90.9%; Score 179; DB 4; Length 419;
 XX Best Local Similarity 79.5%; Pred. No. 2.5e-21;
 XX Matches 35; Conservative 0; Mismatches 9; Indels 0; Gaps 0;
 OY 1 ANSFLXLRQGSLLRXCICXXICDFXAXKXIFEDVDDTLAFWSKH 44
 DB 1 ANSFLXLRQGSLLRXCICXXICDFXAXKXIFEDVDDTLAFWSKH 44
 XX
 XX RESULT 4
 XX AAE08630
 XX ID AAE08630 standard; protein; 419 AA.
 XX AC AAE08630;
 XX DT 01-NOV-2001 (first entry)
 XX DE Human protein C derivative #4.
 XX KW Human; protein C derivative; anticoagulation activity; thrombosis;
 XX serpin inactivation; acute coronary syndrome; myocardial infarction;
 XX vascular occlusive disorder; hypercoagulable state; angina; sepsis;
 XX disseminated intravascular coagulation; DIC; burn; transplantation;
 KW sickle cell disease; viral haemorrhagic fever; protein C deficiency;
 KW haemolytic uremic syndrome; acute arterial thrombotic occlusion;
 KW thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.
 XX
 OS Homo sapiens.
 XX

FN WO200159084-A1.
 XX 16-AUG-2001.
 PD
 XX
 PF 02-FEB-2001; 2001WO-US001221.
 XX
 PR 11-FEB-2000; 2000US-0181948P.
 XX 14-MAR-2000; 2000US-0189199P.
 XX
 PA (ELIL) LILLY & CO ELI.
 XX
 PI Gerlitz BE, Grinnell BW, Jones BE,
 XX
 DR WPI; 2001-5146G2/56.
 DR N-PSDB; AAD15228.
 XX
 PT Protein C derivative for treating acute coronary syndromes, vascular
 PT occlusive disorders, thrombotic disorders and sepsis, comprises
 PT substitutions at specified amino acid positions.
 XX
 PS Claim 6; Page 50-51; 59pp; English.
 XX
 CC This invention relates to human protein C derivatives and nucleic acid
 CC molecules encoding such derivatives. These derivatives have increased
 CC anticoagulation activity, resistance to serpin inactivation and increased
 CC sensitivity to thrombin activation compared to wild type protein C, and
 CC retains the biological activity of the wild type human protein C. Protein
 CC C derivatives are useful in the manufacture of a medicament for the
 CC treatment of acute coronary syndromes e.g. myocardial infarction and
 CC unstable angina; and disease states predisposing to thrombosis; vascular
 CC intravascular coagulation (DIC), burns, transplantations, thalassaemia,
 CC sickle cell disease, viral haemorrhagic fever and haemolytic uremic
 CC syndrome; sepsis in combination with bacterial permeability increasing
 CC protein; thrombotic disorders in combination with an anti-platelet agent;
 CC protein C deficiency; acute arterial thrombotic occlusion,
 CC thromboembolism or stenosis in coronary, cerebral or peripheral arteries
 CC or in vascular grafts in combination with a thrombolytic agent. Nucleic
 CC acid molecules of the invention are useful for treating humans with
 CC genetically predisposed prothrombotic disorders by gene therapy. The
 CC present sequence is human protein C derivative
 XX
 SQ Sequence 419 AA;
 Query Match 90.9%; Score 179; DB 4; Length 419;
 Best Local Similarity 79.5%; Pred. No. 2.5e-21;
 Matches 35; Conservative 0; Mismatches 9; Indels 0; Gaps 0;
 QY 1 ANSFLXXLRQGSIXRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 DB 1 ANSFLXELRQGSLEKIEICDFEAKXIFEDVDTLAFWSKH 44
 RESULT 5
 AAY18301
 ID AAY18301 standard; peptide; 44 AA.
 AC
 XX AAY18301;
 XX
 DT 17-AUG-1999 (first entry)
 XX
 DE Modified GLA domain of vitamin K-dependent protein.
 XX
 KW GLA domain; mutein; vitamin K-dependent protein; clotting disorder;
 XX therapy.
 KW Homo sapiens.
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Misc-difference 1..44
 FT /note= "Xaa= gamma-carboxyglutamic acid, or glutamic
 FT acid"

XX WO9920767-A1.
 XX
 PD 29-APR-1999.
 XX
 PF 20-OCT-1998; 98WO-US022152.
 XX
 PR 23-OCT-1997; 97US-00955636.
 XX
 PA (MINU) UNIV MINNESOTA.
 XX
 PI Neisestuen GL;
 XX
 DR WPI; 1999-288309/24.
 XX
 PT Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid
 PT domain, useful for treating clotting disorders.
 XX
 PS Claim 9; Page 82; 86pp; English.
 XX
 CC This sequence represents a modified GLA (gamma-carboxyglutamic acid)
 CC domain. The invention relates to a vitamin K-dependent polypeptide
 CC comprising a modified GLA domain containing an amino acid substitution
 CC which enhances membrane binding of the modified polypeptide as compared
 CC to the native polypeptide. The polypeptide is used to treat a clotting
 CC disorder by decreasing or increasing clot formation. Modification of the
 CC GLA domain results in a protein which has enhanced membrane binding
 CC affinity as compared to the native protein
 XX
 SQ Sequence 44 AA;
 Query Match 89.3%; Score 176; DB 2; Length 44;
 Best Local Similarity 97.7%; Pred. No. 7.2e-22;
 Matches 43; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ANSFLXXLRQGSIXRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 DB 1 ANSFLXXLRQGSIXRXCIXXICDFXXAKXIFEDVDTLAFWSKH 44
 RESULT 6
 AAB82676
 ID AAB82676 standard; protein; 419 AA.
 XX
 AC AAB82676;
 XX
 DT 15-OCT-2001 (first entry)
 XX
 DE Human protein C derivative (S11G/Q32E/N33D/L194S/T254S).
 XX
 KW Protein C; human; coronary syndrome; thrombosis; angina;
 KW myocardial infarction; vascular occlusive disorder; hypercoagulation;
 KW sepsis; protein C deficiency; occlusion; thromboembolism; stenosis;
 KW antibacterial; immunosuppressive; thrombolytic; cardiant; antianginal;
 KW anticoagulant; therapy; mutant; mutein.
 XX
 OS Homo sapiens.
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Domain 1..45
 FT /note= "Gla domain"
 FT Modified-site 6
 FT /note= "gamma-carboxylated"
 FT Modified-site 7
 FT /note= "gamma-carboxylated"
 FT Misc-difference 11
 FT /note= "Ser in wild-type protein"
 FT Modified-site 14
 FT /note= "gamma-carboxylated"
 FT Modified-site 16
 FT /note= "gamma-carboxylated"
 FT Modified-site 19

FT	Modified-site	/note= "gamma-carboxylated"
FT		20
FT	Modified-site	/note= "gamma-carboxylated"
FT		25
FT	Modified-site	/note= "gamma-carboxylated"
FT		26
FT	Modified-site	/note= "gamma-carboxylated"
FT		29
FT	Misc-difference	/note= "N-glycosylated"
FT		32
FT	Misc-difference	/note= "Gln in wild-type protein"
FT		33
FT	Disulfide-bond	/note= "Asn in wild-type protein"
FT		50. .69
FT	Disulfide-bond	59. .64
FT	Disulfide-bond	80. .89
FT	Disulfide-bond	98. .109
FT	Disulfide-bond	120. .133
FT	Disulfide-bond	141. .277
FT	Cleavage-site	156. .157
FT		/note= "cleavage makes a 2-chain inactive precursor (155- amino acid light chain attached via a disulfide bond to a 282-amino acid heavy chain)"
FT	Peptide	158. .169
FT		/note= "activation peptide; removal activates the 2-chain zymogen"
FT	Cleavage-site	169. .170
FT		/note= "thrombin cleavage site"
FT	Misc-difference	194
FT		/note= "Leu in wild-type protein"
FT	Disulfide-bond	196. .212
FT	Modified-site	248
FT		/note= "N-glycosylated"
FT	Misc-difference	254
FT		/note= "Thr in wild-type protein"
FT	Modified-site	313
FT		/note= "N-glycosylated"
FT	Modified-site	329
FT		/note= "N-glycosylated"
FT	Disulfide-bond	331. .345
FT	Disulfide-bond	356. .384
XX		
PN		W0200157193-A2.
XX		
PD		09-AUG-2001.
XX		
PF		19-JAN-2001; 2001WO-US000020.
XX		
PR		02-FEB-2000; 2000US-0179801P.
PR		14-MAR-2000; 2000US-0189197P.
XX		
PA		(ELIL) LILLY & CO ELI.
XX		
PI		Gerlitz BE, Jones BE;
XX		
DR		WPI; 2001-496919/54.
DR		N-PSDB; AAH26364.
XX		
PT		Novel human protein C derivative for treating, e.g., myocardial infarction, unstable angina, sepsis, thrombotic disorders, acute arterial thrombotic occlusion, and thromboembolism.
XX		
PS		Claim 4; Page 53-54; 63pp; English.
XX		
CC		The present sequence is that of a claimed human protein C derivative in which Ser at position 11 of the mature wild-type protein C sequence (see AAB82673) is substituted with Gly, Gln at position 32 with Glu, Asn at position 33 with Asp, Leu at position 194 with Ser, and Thr at position 254 with Ser. It is an example of protein C derivatives of the invention that have at least 2 amino acid substitutions, but which have increased anticoagulant activity and resistance to inactivation by serpins compared with the wild-type protein, while retaining the biological activity of the wild-type protein. A method of producing the derivatives using

FT Disulfide-bond 120..133
 FT Disulfide-bond 141..277
 FT Cleavage-site 156..157
 /note= "cleavage makes a 2-chain inactive precursor (155-
 FT amino acid light chain attached via a disulfide bond to a
 FT 262-amino acid heavy chain)"
 FT Peptide 158..169
 /note= "activation peptide; removal activates the 2-chain
 FT zymogen"
 FT Cleavage-site 169..170
 /note= "thrombin cleavage site"
 FT Misc-difference 194
 /note= "Leu in wild-type protein"
 FT Disulfide-bond 196..212
 FT Modified-site 248
 /note= "N-glycosylated"
 FT Modified-site 313
 /note= "N-glycosylated"
 FT Modified-site 329
 /note= "N-glycosylated"
 FT Disulfide-bond 331..345
 FT Disulfide-bond 356..384
 WO200157193-A2.
 PN
 XX
 XX
 PD 09-AUG-2001.
 XX
 XX
 PF 19-JAN-2001; 2001WO-US000020.
 XX
 XX
 PR 02-FEB-2000; 2000US-0179801P.
 PR 14-MAR-2000; 2000US-0189197P.
 XX
 XX
 PA (ELIL) LILLY & CO ELI.
 XX
 XX
 PI Gerlitz BE, Jones BE;
 XX
 DR WPI; 2001-496919/54.
 DR N-PSDB; AAH26363.
 XX
 PT Novel human protein C derivative for treating, e.g., myocardial
 PT infarction, unstable angina, sepsis, thrombotic disorders, acute arterial
 FT thrombotic occlusion, and thromboembolism.
 XX
 XX
 PS Claim 3; Page 52-53; 63pp; English.
 XX
 CC The present sequence is that of a claimed human protein C derivative in
 CC which Ser at amino acid position 11 of the mature wild-type protein C
 CC sequence (see AAB82673) is substituted with Gly, Gln at position 32 with
 CC Glu, Asn at position 33 with Asp, and Leu at position 194 with Ser. The
 CC protein is an example of protein C derivatives of the invention that have
 CC at least 2 amino acid substitutions, but which have increased
 CC anticoagulant activity and resistance to inactivation by serpins compared
 CC with the wild-type protein, while retaining the biological activity of
 CC the wild-type protein. A method of producing the derivatives using
 CC recombinant DNA methods is claimed. The protein C derivatives are useful
 CC for treating coronary syndromes and disease states predisposing to
 CC thrombosis (e.g. myocardial infarction and unstable angina), vascular
 CC occlusive disorders and hypercoagulable states, sepsis (in combination
 CC with bactericidal permeability increasing protein or with tissue factor
 CC pathway inhibitor), thrombotic disorders (in combination with an anti-
 CC platelet agent or by local delivery through an intracoronary catheter),
 CC protein C deficiency, acute arterial thrombotic occlusion,
 CC thromboembolism, or stenosis in coronary, cerebral or peripheral arteries
 CC or in vascular grafts. Human patients with genetically predisposed
 CC prothrombotic disorders may be treated by gene therapy (all claimed)
 XX
 XX
 SQ Sequence 419 AA;

Query Match 88.3%; Score 174; DB 4; Length 419;
 Best Local Similarity 77.3%; Pred. No. 1.7e-20;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 QY 1 ANSFLXLRQGSLLXRCIXXCIFDFFXAKXIFEDVDDTLAFWSKH 44

Db 1 ANSFLXLRHGSLEECIEICDFEAKEIFEDVDDTLAFWSKH 44
 RESULT 8
 AAE08628
 ID AAE08628 standard; protein; 419 AA.
 XX
 AC AAE08628;
 XX
 DT 01-NOV-2001 (first entry)
 XX
 DE Human protein C derivative #2.
 XX
 KW Human; protein C derivative; anticoagulation activity; thrombosis;
 KW serpin inactivation; acute coronary syndrome; myocardial infarction;
 KW vascular occlusive disorder; hypercoagulable state; angina; sepsis;
 KW disseminated intravascular coagulation; DIC; burn; transplantation;
 KW sickle cell disease; viral haemorrhagic fever; protein C deficiency;
 KW haemolytic uraemic syndrome; acute arterial thrombotic occlusion;
 KW thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.
 XX
 OS Homo sapiens.
 XX
 PN WO200159084-A1.
 XX
 PD 16-AUG-2001.
 XX
 XX
 PF 02-FEB-2001; 2001WO-US001221.
 XX
 XX
 PR 11-FEB-2000; 2000US-0181948P.
 PR 14-MAR-2000; 2000US-0189199P.
 XX
 XX
 PA (ELIL) LILLY & CO ELI.
 XX
 XX
 PI Gerlitz BE, Grinnell BW, Jones BE;
 XX
 DR WPI; 2001-514662/56.
 DR N-PSDB; AAD15226.
 XX
 PT Protein C derivative for treating acute coronary syndromes, vascular
 PT occlusive disorders, thrombotic disorders and sepsis, comprises
 PT substitutions at specified amino acid positions.
 XX
 PS Claim 4; Page 47-48; 59pp; English.
 XX
 CC The invention relates to human protein C derivatives and nucleic acid
 CC molecules encoding such derivatives. These derivatives have increased
 CC anticoagulation activity, resistance to serpin inactivation and increased
 CC sensitivity to thrombin activation compared to wild type protein C, and
 CC retains the biological activity of the wild type human protein C. Protein
 CC C derivatives are useful in the manufacture of a medicament for the
 CC treatment of acute coronary syndromes e.g. myocardial infarction and
 CC unstable angina; and disease states predisposing to thrombosis; vascular
 CC occlusive disorders and hypercoagulable states e.g. disseminated
 CC intravascular coagulation (DIC), burns, transplantations, thalassaemia,
 CC sickle cell disease, viral haemorrhagic fever and haemolytic uraemic
 CC syndrome; sepsis in combination with bacterial permeability increasing
 CC protein; thrombotic disorders in combination with an anti-platelet agent;
 CC thromboembolism or stenosis in coronary, cerebral or peripheral arteries
 CC or in vascular grafts in combination with a thrombolytic agent. Nucleic
 CC acid molecules of the invention are useful for treating humans with.
 CC genetically predisposed prothrombotic disorders by gene therapy. The
 CC present sequence is human protein C derivative
 XX
 XX
 SQ Sequence 419 AA;

Query Match 88.3%; Score 174; DB 4; Length 419;
 Best Local Similarity 77.3%; Pred. No. 1.7e-20;
 Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
 QY 1 ANSFLXLRQGSLLXRCIXXCIFDFFXAKXIFEDVDDTLAFWSKH 44

Db 1 ANSFLEELRHGSLERECIEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 9
AAE08627
ID AAE08627 standard; protein; 419 AA.

XX AAE08627;

DT 01-NOV-2001 (first entry)

DE Human protein C derivative #1.

XX Human; protein C derivative; anticoagulation activity; thrombosis;
KW serpin inactivation; acute coronary syndrome; myocardial infarction;
KW vascular occlusive disorder; hypercoagulable state; angina; sepsis;
KW disseminated intravascular coagulation; DIC; burn; transplantation;
KW sickle cell disease; viral haemorrhagic fever; protein C deficiency;
KW haemolytic uremic syndrome; acute arterial thrombotic occlusion;
KW thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.

XX Homo sapiens.

XX WO200159084-A1.

XX 16-AUG-2001.

XX 02-FEB-2001; 2001WO-US001221.

XX 11-FEB-2000; 2000US-0181948P.

XX 14-MAR-2000; 2000US-0189199P.

XX (ELIL) LILLY & CO ELI.

XX Gerlitz BE, Grinnell BW, Jones BE;

XX WPI; 2001-514562/56.

XX N-PSDB; AAD15225.

XX Protein C derivative for treating acute coronary syndromes, vascular
PT occlusive disorders, thrombotic disorders and sepsis, comprises
PT substitutions at specified amino acid positions.

XX Claim 3; Page 46-47; 59pp; English.

XX The invention relates to human protein C derivatives and nucleic acid
CC molecules encoding such derivatives. These derivatives have increased
CC anticoagulation activity, resistance to serpin inactivation and increased
CC sensitivity to thrombin activation compared to wild type protein C, and
CC retains the biological activity of the wild type human protein C. Protein
CC C derivatives are useful in the manufacture of a medicament for the
CC treatment of acute coronary syndromes e.g. myocardial infarction and
CC unstable angina; and disease states predisposing to thrombosis; vascular
CC occlusive disorders and hypercoagulable states e.g. disseminated
CC intravascular coagulation (DIC), burns, transplantations, thalassaemia,
CC sickle cell disease, viral haemorrhagic fever and haemolytic uremic
CC syndrome; sepsis in combination with bacterial permeability increasing
CC protein; thrombotic disorders in combination with an anti-platelet agent;
CC protein C deficiency; acute arterial thrombotic occlusion;
CC thromboembolism or stenosis in coronary, cerebral or peripheral arteries
CC or in vascular grafts in combination with a thrombolytic agent. Nucleic
CC acid molecules of the invention are useful for treating humans with
CC genetically predisposed prothrombotic disorders by gene therapy. The
CC present sequence is human protein C derivative

XX Sequence 419 AA;

Query Match 88.3%; Score 174; DB 4; Length 419;
Best Local Similarity 77.3%; Pred. No. 1.7e-20;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

OY 1 ANSFLEXLRQSLRXRCIXXCDFEAKCIFEDVDDTLAFWSKH 44

Query Match 88.3%; Score 174; DB 4; Length 419;

Db 1 ANSFLEELRHGSLERECIEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 10
AAE08629

ID AAE08629 standard; protein; 419 AA.

XX AAE08629;

DT 01-NOV-2001 (first entry)

DE Human protein C derivative #3.

XX Human; protein C derivative; anticoagulation activity; thrombosis;
KW serpin inactivation; acute coronary syndrome; myocardial infarction;
KW vascular occlusive disorder; hypercoagulable state; angina; sepsis;
KW disseminated intravascular coagulation; DIC; burn; transplantation;
KW sickle cell disease; viral haemorrhagic fever; protein C deficiency;
KW haemolytic uremic syndrome; acute arterial thrombotic occlusion;
KW thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.

XX Homo sapiens.

XX WO200159084-A1.

XX 16-AUG-2001.

XX 02-FEB-2001; 2001WO-US001221.

XX 11-FEB-2000; 2000US-0181948P.

XX 14-MAR-2000; 2000US-0189199P.

XX (ELIL) LILLY & CO ELI.

XX Gerlitz BE, Grinnell BW, Jones BE;

XX WPI; 2001-514562/56.

XX N-PSDB; AAD15227.

XX Protein C derivative for treating acute coronary syndromes, vascular
PT occlusive disorders, thrombotic disorders and sepsis, comprises
PT substitutions at specified amino acid positions.

XX Claim 5; Page 48-49; 59pp; English.

XX The invention relates to human protein C derivatives and nucleic acid
CC molecules encoding such derivatives. These derivatives have increased
CC anticoagulation activity, resistance to serpin inactivation and increased
CC sensitivity to thrombin activation compared to wild type protein C, and
CC retains the biological activity of the wild type human protein C. Protein
CC C derivatives are useful in the manufacture of a medicament for the
CC treatment of acute coronary syndromes e.g. myocardial infarction and
CC unstable angina; and disease states predisposing to thrombosis; vascular
CC occlusive disorders and hypercoagulable states e.g. disseminated
CC intravascular coagulation (DIC), burns, transplantations, thalassaemia,
CC sickle cell disease, viral haemorrhagic fever and haemolytic uremic
CC syndrome; sepsis in combination with bacterial permeability increasing
CC protein; thrombotic disorders in combination with an anti-platelet agent;
CC protein C deficiency; acute arterial thrombotic occlusion;
CC thromboembolism or stenosis in coronary, cerebral or peripheral arteries
CC or in vascular grafts in combination with a thrombolytic agent. Nucleic
CC acid molecules of the invention are useful for treating humans with
CC genetically predisposed prothrombotic disorders by gene therapy. The
CC present sequence is human protein C derivative

XX Sequence 419 AA;

Best Local Similarity 77.3%; Pred. No. 1.7e-20;
Matches 34; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 ANSFLXLRQGSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44
DB 1 ANSFLXLRQGSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44

RESULT 11
AAV18298
ID AAV18298 standard; peptide; 44 AA.
AC AAV18298;
XX
XX 17-AUG-1999 (first entry)
DE Modified GLA domain of vitamin K-dependent protein.
XX GLA domain; mutein; vitamin K-dependent protein; clotting disorder;
KW therapy.
XX
XX Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
FT Misc-difference 1. .44 /note= "Xaa= gamma-carboxyglutamic acid, or glutamic
FT acid"
XX
XX WO9920767-A1.
PN
XX 29-APR-1999.
PD
XX 20-OCT-1998; 98WO-US022152.
PF
XX 23-OCT-1997; 97US-00955636.
PR
XX (MINU) UNIV MINNESOTA.
PA
XX Nelsestuen GL;
PI
XX WPI; 1999-288309/24.
PN
XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid
XX domain, useful for treating clotting disorders.
XX
XX Claim 7; Page 78; 86pp; English.
PS
XX This sequence represents a modified GLA (gamma-carboxyglutamic acid)
XX domain. The invention relates to a vitamin K-dependent polypeptide
XX comprising a modified GLA domain containing an amino acid substitution
XX which enhances membrane binding of the modified polypeptide as compared
XX to the native polypeptide. The polypeptide is used to treat a clotting
XX disorder by decreasing or increasing clot formation. Modification of the
XX GLA domain results in a protein which has enhanced membrane binding
XX affinity as compared to the native protein
XX
XX Sequence 44 AA;
SQ

Query Match 87.8%; Score 173; DB 2; Length 44;
Best Local Similarity 97.7%; Pred. No. 2.3e-21;
Matches 43; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANSFLXLRQGSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44
DB 1 ANSFLXLRQGSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44

RESULT 12
AAV18299
ID AAV18299 standard; peptide; 44 AA.
XX
XX AAV18299;
AC

XX 17-AUG-1999 (first entry)
DT Modified GLA domain of vitamin K-dependent protein.
XX GLA domain; mutein; vitamin K-dependent protein; clotting disorder;
XX therapy.
KW
XX
XX Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
FT Misc-difference 1. .44 /note= "Xaa= gamma-carboxyglutamic acid, or glutamic
FT acid"
XX
XX WO9920767-A1.
PN
XX 29-APR-1999.
PD
XX 20-OCT-1998; 98WO-US022152.
PF
XX 23-OCT-1997; 97US-00955636.
PR
XX (MINU) UNIV MINNESOTA.
PA
XX Nelsestuen GL;
PI
XX WPI; 1999-288309/24.
PN
XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic acid
XX domain, useful for treating clotting disorders.
XX
XX Claim 8; Page 78; 86pp; English.
PS
XX This sequence represents a modified GLA (gamma-carboxyglutamic acid)
XX domain. The invention relates to a vitamin K-dependent polypeptide
XX comprising a modified GLA domain containing an amino acid substitution
XX which enhances membrane binding of the modified polypeptide as compared
XX to the native polypeptide. The polypeptide is used to treat a clotting
XX disorder by decreasing or increasing clot formation. Modification of the
XX GLA domain results in a protein which has enhanced membrane binding
XX affinity as compared to the native protein
XX
XX Sequence 44 AA;
SQ

Query Match 86.3%; Score 170; DB 2; Length 44;
Best Local Similarity 95.5%; Pred. No. 7.4e-21;
Matches 42; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANSFLXLRQGSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44
DB 1 ANSFLXLRQGSILRXICIXXICDFXXAKXIFEDVDDTLAFWSKH 44

RESULT 13
AAV18297
ID AAV18297 standard; peptide; 44 AA.
XX
XX AAV18297;
AC
XX 17-AUG-1999 (first entry)
DT
XX Modified GLA domain of vitamin K-dependent protein.
DE GLA domain; mutein; vitamin K-dependent protein; clotting disorder;
XX therapy.
KW
XX Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
FT Misc-difference 1. .44

PS Example 1; Page; 50pp; English.

XX The present sequence is the protein sequence of a mutated Gla domain (N-
CC terminal amino acids 1-45) of human protein C. The mutated Gla domain
CC contains the substitution mutations H10Q, S11G and S12N. Protein C and
CC activated protein C variants comprising a mutated Gla domain are provided
CC by the invention. The variants contain at least 5, and optionally 7-10,
CC amino acid substitutions. A preferred mutant (designated QGNSEEDY, see
CC ABB79946) has the mutations H10Q, S11G, S12N, D23S, Q32E, N33D and H44Y,
CC and shows greatly enhanced anticoagulant activity in standard in vitro
CC coagulation assays. The present mutant (designated QGN) was produced in
CC an example from the invention as a step toward the production of the
CC QGNSEEDY mutant Gla domain, and shows little, if any, improvement in
CC anticoagulant activity over wild-type activated protein C. The invention
CC provides methods for producing the variants based on DNA technology, and
CC with the use of the variants for the treatment of coagulation disorders
CC such as thrombosis or APC resistance, or in diagnostic test systems for
CC assaying components of the protein C-anticoagulant system (all claimed).
CC The variants may also be used in treating arteriosclerosis, myocardial
CC infarction, and disseminated intravascular coagulation. Note: The present
CC sequence is not shown in the specification but is derived from the human
CC wild-type Gla domain sequence given on page 7 of the specification (see
CC ABB79947)

XX SQ Sequence 45 AA;

Query Match 85.3%; Score 168; DB 5; Length 45;
Best Local Similarity 72.7%; Pred. No. 1.6e-20;
Matches 32; Conservative 3; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXLRQGSLLRXRCIXXICDFXXAXXIFEDVDDTLAFWSKH 44
DB 1 ANSFLXLRQGSLLRXRCIXXICDFXXAXXIFEDVDDTLAFWSKH 44

Search completed: March 1, 2004, 10:01:23
Job time : 50 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: March 1, 2004, 09:59:33 ; Search time 13.5 Seconds
(without alignments)
313.513 Million cell updates/sec

Title: SEQ1-4SUBS
Perfect score: 197
Sequence: 1 ANSFLXLRGGSLRXCIXX.....XXAKXIFEDVDTLAFWSKH 44

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283366 seqs, 96191526 residues
Total number of hits satisfying chosen parameters: 283366

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 78.*

1: Piri:.*
2: Piri:.*
3: Piri:.*
4: Piri:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
1	160	81.2	461	1 KXHU	protein C (activat
2	140	71.1	461	1 JX0210	protein C (activat
3	139	70.6	461	1 S18894	protein C (activat
4	122	61.9	456	1 KXHU	coagulation factor
5	115	58.4	482	1 EXBT	coagulation factor
6	114	57.9	492	1 EXBO	coagulation factor
7	110	55.8	488	1 EXHU	coagulation factor
8	101	51.3	443	2 I46932	coagulation factor
9	99	50.3	466	1 KFHU7	coagulation factor
10	86.5	43.9	617	2 S10511	thrombin (EC 3.4.2
11	86.5	43.9	618	2 A35827	coagulation factor
12	86	43.7	475	1 EXCH	coagulation factor
13	85	43.1	407	1 KFB07	plasma protein S p
14	85	43.1	642	2 S53434	plasma protein S p
15	85	43.1	676	1 KXHU5	plasma protein S p
16	84	42.6	622	1 TBHU	thrombin (EC 3.4.2
17	81	41.1	646	2 S38819	plasma protein S -
18	80	40.5	452	1 A30351	coagulation factor
19	80	40.6	459	2 JC0419	coagulation factor
20	80	40.6	461	1 KFHU	coagulation factor
21	80	40.6	675	1 KXBOS	plasma protein S p
22	78	39.6	642	2 S53433	plasma protein S p
23	78	39.6	675	1 KXRTS	plasma protein S p
24	73	37.1	416	1 KFB0	coagulation factor
25	72	36.5	625	1 TBBO	thrombin (EC 3.4.2
26	71	36.0	675	1 KXMS	plasma protein S p
27	59.5	35.3	396	1 KXBOZ	plasma protein 2 -
28	55.5	33.2	422	1 KXHUZ	plasma protein 2 p
29	65	33.0	673	2 A48069	growth arrest-spec

RESULT 1

KXHU

Protein C (activated) (EC 3.4.21.69) precursor - human
N:Alternate names: autoprothrombin IIa; plasma protein C

C:Species: Homo sapiens (man)
C:Date: 17-Mar-1987 #sequence revision 17-Mar-1987 #text_change 16-Jul-1999
C:Accession: A22331; A25426; A21781; A23789; A00927
R:Foster, D.C.; Yoshitake, S.; Davie, E.W.

Proc. Natl. Acad. Sci. U.S.A. 82, 4673-4677, 1985

A:Title: The nucleotide sequence of the gene for human protein C.

A:Reference number: A22331; MUID:85270390; PMID:2991887

A:Accession: A22331

A:Molecule type: DNA

A:Residues: 1-461 <FOS1>

A:Cross-references: GB:M11228; NID:gl90333; PIDN:AAA60166.1; PID:gl90334

R:Plutsky, J.; Hoskins, J.A.; Long, G.L.; Crabtree, G.R.

Proc. Natl. Acad. Sci. U.S.A. 83, 546-550, 1986

A:Title: Evolution and organization of the human protein C gene.

A:Reference number: A25426; MUID:86120978; PMID:3511471

A:Accession: A25426

A:Molecule type: DNA

A:Residues: 1-445, 'L', 446-461 <PLJ>

A:Cross-references: GB:M12712; NID:gl90330; PIDN:AAA60165.1; PID:gl90332

R:Foster, D.; Davie, E.W.

Proc. Natl. Acad. Sci. U.S.A. 81, 4766-4770, 1984

A:Title: Characterization of a cDNA coding for human protein C.

A:Reference number: A21781; MUID:84272714; PMID:6589623

A:Accession: A21781

A:Molecule type: mRNA

A:Residues: 1-107-461 <FOS2>

A:Cross-references: GB:K02059; NID:gl90322; PIDN:AAA60164.1; PID:gl90323

R:Beckmann, R.J.; Schmidt, R.J.; Santerre, R.F.; Plutsky, J.; Crabtree, G.R.; Long, G.L.

Nucleic Acids Res. 13, 5233-5247, 1985

A:Title: The structure and evolution of a 461 amino acid human protein C precursor and i

A:Reference number: A23789; MUID:85269639; PMID:2991859

A:Accession: A23789

A:Molecule type: mRNA

A:Residues: 1-461 <BEC>

A:Cross-references: GB:K02750; NID:G35689; PIDN:CAA36528.1; PID:g763120

R:Miletich, J.P.; Broze Jr., G.J.

J. Biol. Chem. 265, 11397-11404, 1990

A:Title: Beta protein C is not glycosylated at asparagine 329. The rate of translation m

A:Reference number: A44605; MUID:90293094; PMID:1694179

A:Contents: annotation; carbohydrate binding sites; activation peptide

A:Note: The alpha form of protein C is glycosylated at Asn-329, and the beta form is not

R:Harris, R.J.; Ling, V.T.; Spellman, M.W.

J. Biol. Chem. 267, 5102-5107, 1992

A:Title: O-linked fucose is present in the first epidermal growth factor domain of facto:

A:Reference number: A44606; MUID:92184750; PMID:1544894

A:Contents: annotation; beta-hydroxyaspartic acid

C:Comment: Protein C is the zymogen of the vitamin K-dependent serine proteinase that in

ivation of factor Va is strongly enhanced by complexing with protein S. Protein C also fi

growth potentiatin
growth arrest-spec
probable MAP kinas
probable MAP kinas
probable MAP kinas
hypothetical prote
hypothetical prote
protein-tyrosine k
hypothetical prote
ammonium transport
mitogen-activated
VSG expression sit
protein-tyrosine k
platelet-derived g
tyrosine kinase re
type II site-speci

ALIGNMENTS

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;233,299,402/Active site: His, Asp, Ser #status predicted

```

fernlund, P.; Stenflo, J.

J. Biol. Chem. 257, 12170-12179, 1982
A:Title: Amino acid sequence of the light chain of bovine protein C.
A:Reference number: A18385; MUID:83007325; PMID:6896876
A:Accession: A18385
A:Molecule type: protein
A:Residues: 40-194 <PER>
A:Note: 82-Lys was also found
R:Brakenberg, T.; Fernlund, P.; Roepstorff, P.; Stenflo, J.
Proc. Natl. Acad. Sci. U.S.A. 80, 1802-1806, 1983
A:Title: beta-Hydroxyaspartic acid in vitamin K-dependent protein C.
A:Reference number: A19316; MUID:83169769; PMID:6572939
A:Contents: annotation; revision to residue 110
R:Stenflo, J.; Fernlund, P.
J. Biol. Chem. 257, 12180-12190, 1982
A:Title: Amino acid sequence of the heavy chain of bovine protein C.
A:Reference number: A18386; MUID:83007326; PMID:6896877
A:Accession: A18386
A:Molecule type: protein
A:Residues: 197-454, 'pv' <STE>
R:Esmon, N.L.; DeBault, L.B.; Esmon, C.T.
J. Biol. Chem. 258, 5548-5553, 1983
A:Title: Proteolytic formation and properties of gamma-carboxyglutamic acid-domainless p
A:Reference number: A37541; MUID:83213513; PMID:6304092
A:Contents: annotation; activation; calcium binding
R:Johnson, A.E.; Esmon, N.L.; Laue, T.M.; Esmon, C.T.
J. Biol. Chem. 258, 5554-5560, 1983
A:Title: Structural changes required for activation of protein C are induced by Ca2+ bin
A:Reference number: A37542; MUID:83213514; PMID:6406503
A:Contents: annotation; activation; calcium binding
C:Comment: Protein C is the zymogen of the vitamin K-dependent serine proteinase that re
s.
C:Comment: Protein C is synthesized in the liver as a single chain precursor, which is c
bin, which cleaves a tetradecapeptide from the amino end of the heavy chain; this reacti
C:Comment: Calcium binds to the gamma-carboxyglutamic acid (Gla) residues and, with stro
cognition of the thrombin-thrombomodulin complex.
C:Comment: The gamma-carboxyglutamic acid residues arise by a posttranslational, vitamin
C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
C:Keywords: anticoagulant; beta-hydroxyaspartic acid; blood coagulation; calcium binding
F:1-29/Domain: signal sequence (fragment) #status predicted <SIG>
F:24-83/Domain: Gla domain homology <GUA>
F:30-39/Domain: propeptide #status predicted <PRO>
F:40-194/Product: protein C light chain #status experimental <LCH>
F:137-172/Domain: EGF homology <EG1>
F:197-456/Product: protein C heavy chain #status experimental <HCH>
F:211-440/Domain: activation peptide #status experimental <APT>
F:45,46,53,55,58,59,62,64,65,68,74/Modified site: gamma-carboxyglutamic acid (Glu) #stat
F:110/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status experimental
F:119-128,137-148,144-157,159-172,180-318,237-253,368-382,393-421/Disulfide bonds: #stat
F:136,289,350/Binding site: carboxylate (Asn) (covalent) #status predicted
F:252,298,397/Active site: His, Asp, Ser #status predicted
F:366/Binding site: carboxylate (Asn) (covalent) #status predicted
Query March 61.9%; Score 122; DB 1; Length 456;
Best Local Similarity 50.0%; Pred. No. 1.2e-11;
Matches 21; Conservative 9; Mismatches 12; Indels 0; Gaps 0;
QY 1 ANSFLXLRQSLXKXCIXICDFXXKXIFEDVDDTLAFWS 42
DB 40 ANSFLXLRQSLXKXCIXICDFXXKXIFEDVDDTLAFWS 81
RESULT 5
EXRT
coagulation factor Xa (EC 3.4.21.6) precursor - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 31-Jan-1995 #sequence revision 07-Feb-1997 #text_change 08-Dec-2000
C:Accession: S49075; JC4670; PS0190; PS0190; PS0190; PS0190; PS0190; PS0190; PS0190; PS0190
R:Stanton, C.; Ross, R.P.; Hutson, S.; Wallin, R.
Thromb. Res. 80, 63-73, 1995
A:Title: Evidence for competition between vitamin K-dependent clotting factors for intra
A:Reference number: A58498; MUID:96093366; PMID:8578539

A:Accession: S49075
A:Molecule type: mRNA
A:Residues: 1-482 <STA1>
A:Cross-references: EMBL:X79807; NID:9506600; PIDN:CAA56202.1; PID:9506601
A:Note: submitted to the EMBL Data Library, June 1994
A:Note: neither the complete nucleic acid sequence nor the complete translation are show
R:Stanton, C.; Ross, R.P.; Hutson, S.; Wallin, R.
Gene 169, 269-273, 1996
A:Title: Processing and expression of rat and human clotting factor X-encoding cDNAs.
A:Reference number: JC4670; MUID:96194815; PMID:8647460
A:Accession: JC4670
A:Molecule type: mRNA
A:Residues: 1-482 <STA2>
A:Cross-references: EMBL:X79807; NID:9506600; PIDN:CAA56202.1; PID:9506601
A:Experimental source: Cos-1 cell
R:Enyoji, K.; Miyazaki, K.; Kato, H.
J. Biochem. 109, 890-898, 1991
A:Title: Characterization of rat factors X and Xa: demonstration of factor Xa in rat pla
A:Reference number: PS0190; MUID:92041742; PMID:1718949
A:Accession: PS0191
A:Molecule type: protein
A:Residues: 41-58 'X', 60-65 <ENJ1>
A:Accession: PS0190
A:Molecule type: protein
A:Residues: 183-186 'X', 188-207 <ENJ2>
R:Murakawa, M.; Okamura, T.; Kamura, T.; Kuroiwa, M.; Hazada, M.; Niho, Y.
Eur. J. Haematol. 52, 162-168, 1994
A:Title: Analysis of the partial nucleotide sequences and deduced primary structures of
A:Reference number: I46196; MUID:94222160; PMID:8168596
A:Accession: I62745
A:Status: preliminary; translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 295-383 'G', 385-455 <MUR>
A:Cross-references: GB:D21215; NID:9415309; PIDN:BA04756.1; PID:9455396
C:Function:
A:Description: catalyzes the proteolytic activation of prothrombin to thrombin in the pr
A:Pathway: blood coagulation
C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglutam
F:1-23/Domain: signal sequence #status predicted <SIG>
F:24-40/Domain: propeptide #status predicted <PRO>
F:25-84/Domain: Gla domain homology <GUA>
F:41-179/Product: coagulation factor X light chain #status predicted <LCH>
F:90-121/Domain: EGF homology <EG1>
F:129-164/Domain: EGF homology <EG2>
F:183-482/Product: coagulation factor X heavy chain #status predicted <HCH>
F:183-231/Domain: activation peptide #status predicted <APT>
F:232-460/Domain: trypsin homology <TRY>
F:45,47,54,56,59,60,65,66,69,72,79/Modified site: gamma-carboxyglutamic acid (Glu) #stat
F:57-62,90-101,95-110,112-121,129-140,136-149,151-164,172-340,238-243,259-275,388-402,411
F:103/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status predicted
F:187/Binding site: carboxylate (Asn) (covalent) #status experimental
F:208/Binding site: carboxylate (Thr) (covalent) #status predicted
F:218/Binding site: carboxylate (Asn) (covalent) #status predicted
F:231-232/Cleavage site: Arg-Tle (coagulation factor IXa, coagulation factor VIIa) #stat
F:274,320,417/Active site: His, Asp, Ser #status predicted
Query Match 58.4%; Score 115; DB 1; Length 482;
Best Local Similarity 43.2%; Pred. No. 1.6e-10;
Matches 19; Conservative 10; Mismatches 15; Indels 0; Gaps 0;
QY 1 ANSFLXLRQSLXKXCIXICDFXXKXIFEDVDDTLAFWSKH 44
DB 41 ANSFLXLRQSLXKXCIXICDFXXKXIFEDVDDTLAFWSKH 84
RESULT 6
EXBO
coagulation factor Xa (EC 3.4.21.6) precursor - bovine
N:Alternate names: Stuart factor
C:Species: Bos primigenius taurus (cattle)
C:Date: 24-Apr-1984 #sequence_revision 17-Mar-1987 #text_change 16-Jul-1999

C/Accession: A22867; A14997; A12030; A34412; S39414; A00925
R/Fung, M.R.; Campbell, R.M.; MacGillivray, T.A.
Nucleic Acids Res. 12, 4481-4492, 1984
A/Title: Blood coagulation factor X mRNA encodes a single polypeptide chain containing a
A/Reference number: A22867; MUID:84247315; PMID:6330671
A/Accession: A22867
A/Molecule type: mRNA
A/Residues: 1-487 <FUN>
A/Cross-references: GB:X00673; NID:G192; PIDN:CAA25286.1; PID:G193
R/Enfield, D.L.; Ericsson, L.H.; Fujikawa, K.; Walsh, K.A.; Neurath, H.; Titani, K.
Biochemistry 19, 659-667, 1980
A/Title: Amino acid sequence of the light chain of bovine factor X-1 (Stuart factor).
A/Reference number: A14997; MUID:80130563; PMID:6766735
A/Accession: A14997
A/Molecule type: protein
A/Residues: 41-102, 'N', 104-180 <ENF>
R/McMullen, B.A.; Fujikawa, K.; Kisiel, W.
Biochem. Biophys. Res. Commun. 115, 8-14, 1983
A/Title: The occurrence of beta-hydroxyaspartic acid in the vitamin K-dependent blood coagulation factors.
A/Reference number: A20274; MUID:8330813; PMID:6688526
A/Contents: annotation; revision to residue 103
R/Titani, K.; Fujikawa, K.; Enfield, D.L.; Ericsson, L.H.; Walsh, K.A.; Neurath, H.
Proc. Natl. Acad. Sci. U.S.A. 72, 3082-3086, 1975
A/Title: Bovine factor X-1 (Stuart factor): amino-acid sequence of heavy chain.
A/Reference number: A12030; MUID:76053069; PMID:1059093
A/Accession: A12030
A/Molecule type: Protein
A/Residues: 183-292,294-295, 'GDE', 299-334,336-348, 'AE', 351-354,356-441, 'GKFG', 446-492 <T>
A/Note: carboxylate binding sites and disulfide bonds were determined
R/Person, E.; Selander, M.; Linse, S.; Drakenberg, T.; Oehlin, A.K.; Stenflo, J.
J. Biol. Chem. 264, 16897-16904, 1989
A/Title: Calcium binding to the isolated beta-hydroxyaspartic acid-containing epidermal
A/Reference number: A34412; MUID:89380326; PMID:2789221
A/Accession: A34412
A/Molecule type: Protein
A/Residues: 85-126 <PER>
A/Note: beta-hydroxyaspartic acid site
R/Inoue, K.; Morita, T.
Eur. J. Biochem. 218, 153-163, 1993
A/Title: Identification of O-linked oligosaccharide chains in the activation peptides of
A/Reference number: S39414; MUID:94062825; PMID:8243461
A/Accession: S39414
A/Molecule type: protein
A/Residues: 183-196,199-209,216-233 <INO>
A/Note: carboxylate binding sites
R/Titani, K.; Hermanson, M.A.; Fujikawa, K.; Ericsson, L.H.; Walsh, K.A.; Neurath, H.; D
Biochemistry 11, 4899-4903, 1972
A/Title: Bovine factor X-1a (activated Stuart factor). Evidence of homology with mammal
A/Reference number: A12453; MUID:73053314; PMID:4264286
A/Contents: annotation; active site
R/Fujikawa, K.; Titani, K.; Davie, E.W.
Proc. Natl. Acad. Sci. U.S.A. 72, 3359-3363, 1975
A/Title: Activation of bovine factor X (Stuart factor): conversion of factor Xaalpha to
A/Reference number: A13504; MUID:76053121; PMID:1059122
A/Contents: annotation; activation
R/Sugo, T.; Bjork, I.; Holmgren, A.; Stenflo, J.
J. Biol. Chem. 259, 5705-5710, 1984
A/Title: Calcium-binding properties of bovine factor X lacking the gamma-carboxyglutamic
A/Reference number: A38024; MUID:84185716; PMID:6546930
A/Contents: annotation; calcium binding
R/Morita, T.; Jackson, C.M.
J. Biol. Chem. 261, 4008-4014, 1986
A/Reference number: A38025; MUID:86440210; PMID:3949800
A/Contents: annotation; sulfate binding
C/Comment: Factor Xa converts prothrombin to thrombin during blood clotting.
C/Comment: The two chains are formed from a single-chain precursor by the excision of two
C/Comment: The activation peptide is cleaved by factor IXa (in the intrinsic pathway), C
activation.
C/Comment: Calcium binds to the gamma-carboxyglutamic acid (Gla) residues and, with str
C/Comment: The gamma-carboxyglutamic acid residues arise by a posttranslational, vitamin
C/Genetics:
A/Genes: F10
A/Map position: 13q34

C/Function:
A/Description: catalyzes the proteolytic activation of prothrombin to thrombin in the pr
A/Pathway: blood coagulation
C/Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
C/Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglutam
F1-15/Domain: signal sequence #status predicted <SIG>
F16-40/Domain: propeptide #status predicted <PRO>
F25-84/Domain: Gla domain homology <GUA>
F25-84/Product: coagulation factor X light chain #status experimental <LCH>
F90-120/Domain: EGF homology <EG1>
F183-492/Domain: EGF homology <EG2>
F183-492/Product: coagulation factor X heavy chain #status experimental <HCH>
F183-233/Domain: activation peptide #status experimental <APT>
F234-492/Product: coagulation factor Xa heavy chain #status experimental <AHC>
F234-461/Domain: trypsin homology <TRI>
F46,47,54,56,59,60,65,66,69,72,75,79/Modified site: gamma-carboxyglutamic acid (Glu) #s
F57-62,90-101,95-110,112-121,129-140,136-149,151-164,172-341/Disulfide bonds: #status p
F103/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status experimental
F200/Binding site: sulfate (Tyr) (covalent) (partial) #status experimental
F208,485/Binding site: carbohydrate (Thr) (covalent) #status experimental
F218/Binding site: carbohydrate (Asn) (covalent) #status experimental
F233-234/Cleavage site: Arg-116 (coagulation factor IXa, coagulation factor VIIa) #stat
F240-245,250-278,389-403,414-442/Disulfide bonds: #status experimental
F275,321,418/Active site: His, Asp, Ser #status predicted
Query Match 57.9%; Score 114; DB 1; Length 492;
Best Local Similarity 45.5%; Pred. No. 2.4e-10;
Matches 20; Conservative 8; Mismatches 16; Indels 0; Gaps 0;
QY 1 ANSTLXLROSLRXKICIXICDFKXAXKIFEDVDDTLAFWSKH 44
Db 41 ANSFLKVKQGNRECLLEACSLSEAREVFEDEAQTDFWSKY 84
RESULT 7
EXHU
coagulation factor Xa (EC 3.4.21.6) precursor [validated] - human
N/Alternate names: Stuart factor
C/Species: Homo sapiens (man)
C/Date: 15-Nov-1984 #sequence revision 02-May-1994 #text change 08-Dec-2000
A/Accession: A24478; JQ0917; A42485; A25853; A22208; A21284; A20362; S39415; I54051; A00
R/Leytus, S.P.; Foster, D.C.; Kurachi, K.; Davie, E.W.
Biochemistry 25, 5098-5102, 1986
A/Title: Gene for human Factor X: a blood coagulation factor whose gene organization is
A/Reference number: A24478; MUID:87026500; PMID:3768336
A/Accession: A24478
A/Molecule type: DNA
A/Residues: 1-488 <LBY>
A/Cross-references: GB:L29433; GB:M14327; NID:9459809; PIDN:AAA52764.1; PID:G182831
R/Messier, T.L.; Pittman, D.D.; Long, G.L.; Kaufman, R.J.; Church, W.R.
Gene 99, 291-294, 1991
A/Title: Cloning and expression in COS-1 cells of a full-length cDNA encoding human coag
A/Reference number: JQ0917; MUID:91216473; PMID:1902434
A/Accession: JQ0917
A/Molecule type: mRNA
A/Residues: 1-488 <MES>
R/Miao, C.H.; Leytus, S.P.; Chung, D.W.; Davie, E.W.
J. Biol. Chem. 267, 7395-7401, 1992
A/Title: Liver-specific expression of the gene coding for human factor X, a blood coagul
A/Reference number: A42485; MUID:92218390; PMID:1313796
A/Accession: A42485
A/Molecule type: DNA
A/Residues: 1-15 <MIA>
A/Experimental source: liver
A/Note: sequence extracted from NCBI backbone (NCBI:93780, NCBI:93787)
R/Kaul, R.K.; Hildebrand, B.; Roberts, S.; Jagadeeswaran, P.
Gene 41, 311-314, 1986
A/Title: Isolation and characterization of human blood-coagulation factor X cDNA.
A/Reference number: A25853; MUID:86221713; PMID:3011603
A/Accession: A25853
A/Molecule type: mRNA
A/Residues: 19-284, 'E', 289-488 <KAU>

RESULT 9
KFHUT
coagulation factor VIIa (EC 3.4.21.21) precursor [validated] - human
C.Species: Homo sapiens (man)
C.Date: 19-May-1989 sequence_revision 19-May-1994 #text_change 08-Dec-2000
C.Accession: A28322; A23819; A31186; B31186; S63524
C.Prov.Hara, P.J.; Grant, F.J.; Haldeman, B.A.; Gray, C.L.; Inasley, M.Y.; Hagen, P.S.; Murr
Proc. Natl. Acad. Sci. U.S.A. 84, 5158-5162, 1987
A.Title: Nucleotide sequence of the gene coding for human factor VII, a vitamin K-depend
A.Reference number: A28322; MUID:87260949; PMID:3037537
A.Accession: A28322
A.Molecule type: DNA
A.Residues: 1-466 <OHA>
A.Cross-references: GB:J02933; NID:gi80333; PIDN:AAA51983.1; PID:gi180334
R:Hagen, P.S.; Gray, C.L.; O'Hara, P.; Grant, F.J.; Saari, G.C.; Woodbury, R.G.; Hart, C
Proc. Natl. Acad. Sci. U.S.A. 83, 2412-2416, 1986
A.Title: Characterization of a cDNA coding for human factor VII.
A.Reference number: A23819; MUID:86205965; PMID:3486420
A.Accession: A23819
A.Molecule type: mRNA
A.Residues: 1-466 <HAG>
A.Cross-references: GB:M13232; NID:gi182799; PIDN:AAA88040.1; PID:gi182801
R:Thim, L.; Bjoern, S.; Christensen, M.; Nicolaisen, E.M.; Lund-Hansen, T.; Pedersen, A.J
Biochemistry 27, 7785-7793, 1988
A.Title: Amino acid sequence and posttranslational modifications of human factor VII-a f
A.Reference number: A90539; MUID:89088153; PMID:3264725
A.Accession: A31186
A.Molecule type: protein

A:Residues: 61-212 <THR>
 A:Accession: B31186
 A:Molecule type: protein
 A:Residues: 213-466 <THR>
 R:Bjorn, S.; Foster, D.C.; Thim, L.; Wiberg, F.C.; Christensen, M.; Komiyama, Y.; Peder
 J. Biol. Chem. 266, 11051-11057, 1991
 A:Title: Human plasma and recombinant factor VII. Characterization of O-glycosylations a
 A:Reference number: A4059; MUID:91250411; PMID:1904059
 A:Contents: annotation; carbohydrate binding sites
 R:Persson, E.; Petersen, L.C.
 Eur. J. Biochem. 234, 293-300, 1995
 A:Title: Structurally and functionally distinct Ca(2+) binding sites in the gamma-carbox
 A:Reference number: S63524; MUID:96096752; PMID:8529655
 A:Accession: S63524
 A:Molecule type: protein
 A:Residues: 61-65;99-103;105-109;213-217;308-312 <PER>
 C:Genetics:
 A:Gens: GDB:F7
 A:Cross-references: GDB:119897; OMIM:227500
 A:Map position: 13q34-13q34
 A:Introns: 22/1; 44/1; 97/3; 106/1; 144/1; 191/1; 227/3; 269/1
 C:Function:
 A:Description: catalyzes the proteolytic activation of coagulation factor X in the prese
 coagulation factor IX in the presence of calcium and tissue factor
 A:Pathway: blood coagulation extrinsic pathway
 C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology
 C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglutam
 F:1-20/Domain: signal sequence #status predicted <SIG>
 F:21-60/Domain: propeptide #status predicted <PRO>
 F:45-104/Domain: Gla domain homology <GLA>
 F:61-212/Product: coagulation factor VIIa light chain #status experimental <MA1>
 F:110-141/Domain: EGF homology <EG1>
 F:151-187/Domain: EGF homology <EG2>
 F:213-466/Product: coagulation factor VIIa heavy chain #status experimental <MA2>
 F:213-447/Domain: trypsin homology <TRY>
 F:66-67;74;76;79;80;85;86;89;95/Modified site: gamma-carboxyglutamic acid (Glu) #status
 F:77-82;110-121;115-130;132-141;151-162;158-172;174-187;195-322;219-224;238-254;370-389,
 F:112;120/Binding site: carboxyhydrate (Ser) (covalent) #status experimental
 F:123/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status absent
 F:205;382/Binding site: carboxyhydrate (Asn) (covalent) #status experimental
 F:212-213/Cleavage site: Arg-His (coagulation factor X) #status experimental
 F:253;302;404/Active site: His, Asp, Ser #status predicted
 F:350-351/Cleavage site: Arg-Gly (coagulation factor Xa) #status predicted

Query Match 50.3%; Score 99; DB 1; Length 466;
 Best Local Similarity 48.8%; Pred. No. 6;1e-08;
 Matches 20; Conservative 4; Mismatches 17; Indels 0; Gaps 0;

QY 1 ANSFLXLRQSLRXPXIXXICDFXXAKXIFEDVDDTLAFW 41
 DB 61 ANAFLELRPGSLRECKEQCSFEAREIFPKDAERTKLEW 101

RESULT 10
 S10511
 thrombin (EC 3.4.21.5) precursor - rat
 C:Species: Rattus norvegicus (Norway rat)
 C:Date: 07-May-1993 #sequence revision 07-May-1993 #text_change 03-May-2002
 A:Accession: S10511; A60576; B42696
 R:Diwanich, M.; Monard, D.
 Nucleic Acids Res. 18, 4251, 1990
 A:Title: cDNA sequence of rat prothrombin.
 A:Reference number: S10511; MUID:90332426; PMID:2377469
 A:Accession: S10511
 A:Molecule type: mRNA
 A:Residues: 1-617 <DIH>
 A:Cross-references: EMBL:X52835; NID:956969; PIDN:CAA37017.1; PID:956970
 R:Henrikson, K.P.; Jasin, E.B.; Greenwood, J.A.; Dickerman, H.W.
 Endocrinology 126, 167-175, 1990
 A:Title: Prothrombin levels are increased in the estrogen-treated immature rat uterus.
 A:Reference number: A60576; MUID:90091942; PMID:2233980
 A:Accession: A60576
 A:Molecule type: protein

A:Residues: 44-58 <HEN>
 A:Note: the authors purified the proenzyme from the estrogen-stimulated maturing rat ute
 R:Banfield, D.K.; MacGillivray, R.T.A.
 Proc. Natl. Acad. Sci. U.S.A. 89, 2779-2783, 1992
 A:Title: Partial characterization of vertebrate prothrombin cDNAs: amplification and seq
 A:Reference number: A42696; MUID:92212913; PMID:1557383
 A:Accession: B42696
 A:Status: Preliminary
 A:Molecule type: mRNA
 A:Residues: 383-617, 'E', <BAN>
 A:Cross-references: GB:X81394
 C:Superfamily: thrombin; Gla domain homology; kringle homology; trypsin homology
 C:Keywords: blood coagulation; calcium binding; carboxyglutamic acid; glycoprotein; hydr
 F:1-24/Domain: signal sequence #status predicted <SIG>
 F:25-43/Domain: propeptide #status predicted <PRO>
 F:28-88/Domain: Gla domain homology <GLA>
 F:44-617/Product: prothrombin #status experimental <PMAT>
 F:109-187/Domain: kringle homology <KR1>
 F:215-292/Domain: kringle homology <KR2>
 F:360-609/Domain: trypsin homology <TRY>
 F:50;51;58;60;63;64;69;70;73;76/Modified site: gamma-carboxyglutamic acid (Glu) #status
 F:61-66;91-104;109-187;130-170;158-182;215-292;236-276;284-387;332-478;387-403;532-546;5
 F:402;458;564/Active site: His, Asp, Ser #status predicted

Query Match 43.9%; Score 86.5; DB 2; Length 617;
 Best Local Similarity 42.2%; Pred. No. 8;4e-06;
 Matches 19; Conservative 6; Mismatches 19; Indels 1; Gaps 1;

QY 1 ANS-FLXXLRQSLRXPXIXXICDFXXAKXIFEDVDDTLAFWSKH 44
 DB 44 ANSGFLELRKGNLERECVEQCSFEAREFALSPQDQVFWAKY 88

RESULT 11
 A35827
 thrombin (EC 3.4.21.5) precursor - mouse
 C:Species: Mus musculus (house mouse)
 C:Date: 14-Dec-1990 #sequence revision 14-Dec-1990 #text_change 03-May-2002
 A:Accession: A35827; A43696; S12081
 R:Degeen, S.J.F.; Schaefer, L.A.; Jamison, C.S.; Grant, S.G.; Fitzgibbon, J.J.; Pai, J.A.
 DNA Cell Biol. 9, 481-496, 1990
 A:Title: Characterization of the cDNA coding for mouse prothrombin and localization of t
 A:Reference number: A35827; MUID:91025551; PMID:2222810
 A:Accession: A35827
 A:Status: Preliminary
 A:Molecule type: mRNA
 A:Residues: 1-618 <DEG>
 A:Cross-references: GB:X52308; NID:953813; PIDN:CAA36548.1; PID:953814
 A:Experimental source: strain C57BL/6
 A:Note: the data were obtained from females resulting from the cross of M. domesticus an
 R:Banfield, D.K.; MacGillivray, R.T.A.
 Proc. Natl. Acad. Sci. U.S.A. 89, 2779-2783, 1992
 A:Title: Partial characterization of vertebrate prothrombin cDNAs: amplification and seq
 A:Reference number: A42696; MUID:92212913; PMID:1557383
 A:Accession: A42696
 A:Status: Preliminary
 A:Molecule type: mRNA
 A:Residues: 384-618, 'E', <BAN>
 A:Cross-references: GB:X81394
 C:Superfamily: thrombin; Gla domain homology; kringle homology; trypsin homology
 C:Keywords: blood coagulation; calcium binding; carboxyglutamic acid; glycoprotein; hydr
 F:1-24/Domain: signal sequence #status predicted <SIG>
 F:25-43/Domain: propeptide #status predicted <PRO>
 F:28-88/Domain: Gla domain homology <GLA>
 F:44-617/Product: prothrombin B #status predicted <MAT>
 F:109-187/Domain: kringle homology <KR1>
 F:215-293/Domain: kringle homology <KR2>
 F:361-610/Domain: trypsin homology <TRY>
 F:50;51;58;60;63;64;69;70;73;76/Modified site: gamma-carboxyglutamic acid (Glu) #status
 F:61-66;91-104;109-187;130-170;158-182;215-293;236-276;284-388;333-479;388-404;533-547;5
 F:403;459;565/Active site: His, Asp, Ser #status predicted

Query Match 43.9%; Score 86.5; DB 2; Length 618;

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RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko V., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield V.S.N., Krzywinski M.I., Skalska U., Smallus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.,
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [6]
RP SEQUENCE OF 106-461 FROM N.A.
RX MEDLINE=84272714; PubMed=6589623;
RA Foster D.C., Davie E.W., PubMed=6589623;
RT "Characterization of a cDNA coding for human protein C.";
RN Proc. Natl. Acad. Sci. U.S.A. 81:4766-4770(1984).
RN [7]
RP CARBOHYDRATE-LINKAGE SITE ASN-371.
RX MEDLINE=90293094; PubMed=1694179;
RA Miletich J.P., Broze G.J. Jr.;
RT "Beta protein C is not glycosylated at asparagine 329. The rate of
RT translation may influence the frequency of usage at asparagine-X-
RT cysteine sites.";
RL J. Biol. Chem. 265:11397-11404(1990).
RN [8]
RP HYDROXYLATION.
RX MEDLINE=92184750; PubMed=1544894;
RA Harris R.J., Ling V.T., Spellman M.W.;
RT "O-linked fucose is present in the first epidermal growth factor
RT domain of factor XII but not protein C.";
RL J. Biol. Chem. 267:5102-5107(1992).
RN [9]
RP 3D-STRUCTURE MODELING OF 175-450.
RX MEDLINE=94272342; PubMed=8003977;
RA Fisher C.L., Greengard J.S., Griffin J.H.;
RT "Models of the serine protease domain of the human antithrombotic
RT plasma factor activated protein C and its zymogen.";
RL Plasma Sci. 3:588-599(1994).
RN [10]
RP X-RAY CRYSTALLOGRAPHY (2.8 ANGSTROMS) OF 84-461.
RX MEDLINE=97157472; PubMed=9003757;
RA Mather T., Oganessyan V., Hof P., Huber R., Foundling S., Esmen C.,
RT Bode W.;
RT "The 2.8 A crystal structure of Gla-domainless activated protein C.";
RL EMBO J. 15:6822-6831(1996).
RN [11]
RP REVIEW ON PROC VARIANTS.
RX MEDLINE=93190290; PubMed=8446940;
RA Reitsma P.H., Poort S.R., Bernardi F., Gandrille S., Long G.L.,
RA Sala N., Cooper D.N.;
RT "Protein C deficiency: a database of mutations. For the Protein C & S
RT Subcommittee of the Scientific and Standardization Committee of the
RT International Society on Thrombosis and Haemostasis.";
RL Thromb. Haemost. 69:77-84(1993).
RN [12]
RP VARIANT PROC DEFICIENCY CYS-444.
RX MEDLINE=87204221; PubMed=2437584;
RA Romeo G., Hassan H.J., Scaemphil S., Roncuzzi L., Cianetti L.,
RA Leonardi A., Vicente V., Mannucci P.M., Bertina R.M., Peschle C.,
RA Cortese R.;
RT "Hereditary thrombophilia: identification of nonsense and missense
RT mutations in the protein C gene.";
RL Proc. Natl. Acad. Sci. U.S.A. 84:2829-2832(1987).
RN [13]
RP VARIANT PROC DEFICIENCY TRP-211.
RX MEDLINE=90098906; PubMed=2602169;
RA Grundy C.B., Chitole A., Talbot S., Bevan D., Kakkar V.V.,
RA Cooper D.N.;
RT "Protein C London 1: recurrent mutation at Arg-169 (CGG-->TGG) in
RT the protein C gene causing thrombosis.";
RL Nucleic Acids Res. 17:10513-10513(1989).
RN [14]
RP VARIANT PROC DEFICIENCY CYS-272.
RX MEDLINE=91329836; PubMed=1868249;

RA Reitsma P.H., Poort S.R., Allaart C.F., Briet E., Bertina R.M.;
RT "The spectrum of genetic defects in a panel of 40 Dutch families with
RT symptomatic protein C deficiency type I: heterogeneity and founder
RT effects.";
RL Blood 78:890-894(1991).
RN [15]
RP VARIANTS PROC DEFICIENCY ALA-62 AND MET-76.
RX MEDLINE=92190481; PubMed=1347706;
RA Bovill E.G., Tomczak J.A., Grant B., Bhushan F., Pillemer E.,
RA Rainville I.R., Long G.L.;
RT "Protein Cwerment: symptomatic type II protein C deficiency
RT associated with two GLA domain mutations.";
RL Blood 79:1456-1465(1992).
RN [16]
RP VARIANT PROC DEFICIENCY ASP-418.
RX MEDLINE=92305321; PubMed=1611081;
RA Sugihara Y., Miura O., Yuen P., Aoki N.;
RT "Protein C deficiency Hong Kong 1 and 2: hereditary protein C
RT deficiency caused by two mutant alleles, a 5-nucleotide deletion and
RT a missense mutation.";
RL Blood 80:126-133(1992).
RN [17]
RP VARIANT PROC DEFICIENCY LEU-289.
RX MEDLINE=92380660; PubMed=1511988;
RA Grundy C.B., Chisholm M., Kakkar V.V., Cooper D.N.;
RT "A novel homozygous missense mutation in the protein C (PROC) gene
RT causing recurrent venous thrombosis.";
RL Hum. Genet. 89:683-684(1992).
RN [18]
RP VARIANTS PROC DEFICIENCY GLN-220 AND TRP-220.
RX MEDLINE=92380661; PubMed=1511989;
RA Grundy C.B., Schulman S., Tengborn L., Kakkar V.V., Cooper D.N.;
RT "Two different missense mutations at Arg 178 of the protein C (PROC)
RT gene causing recurrent venous thrombosis.";
RL Hum. Genet. 89:685-686(1992).
RN [19]
RP VARIANT PROC DEFICIENCY GLN-220.
RX MEDLINE=92350852; PubMed=1301959;
RA Gandrille S., Vidau M., Alach M., Alhenc-Gelas M., Fischer A.M.,
RA Gouault-Heilman M., Toulon P., Flessinger J.N., Goossens M.;
RT "Two novel mutations responsible for hereditary type I protein C
RT deficiency: characterization by denaturing gradient gel
RT electrophoresis.";
RL Hum. Mutat. 1:491-500(1992).
RN [20]
RP VARIANT PROC DEFICIENCY SER-334.
RX MEDLINE=92276939; PubMed=1593215;
RA Yamamoto K., Matsushita T., Sugiyama I., Takamatsu J., Iwasaki E.,
RA Wada H., Deguchi K., Shirakawa S., Saito H.;
RT "Homozygous protein C deficiency: identification of a novel missense
RT mutation that causes impaired secretion of the mutant protein C.";
RL J. Lab. Clin. Med. 119:682-689(1992).
RN [21]
RP VARIANTS PROC DEFICIENCY TRP-38; CYS-42; HIS-42; GLN-271 AND ASN-294.
RX MEDLINE=93313192; PubMed=8324221;
RA Gandrille S., Alhenc-Gelas M., Gausem P., Aillaud M.-F., Dupuy E.,
RA Julian-Vague I., Alach M.;
RT "Five novel mutations located in exons III and IX of the protein C
RT gene in patients presenting with defective protein C anticoagulant
RT activity.";
RL Blood 82:159-168(1993).
RN [22]
RP VARIANTS PROC DEFICIENCY GLY-14; GLN-211; TYR-244; GLN-253; LEU-321;
RX CYS-328; ILE-385; THR-388 AND VAL-388.
RX MEDLINE=93271391; PubMed=8499565;
RA Poort S.R., Pabinger-Fasching I., Mannhalter C., Reitsma P.H.,
RA Bertina R.M.;
RT "Twelve novel and two recurrent mutations in 14 Austrian families
RT with hereditary protein C deficiency.";
RL Blood Coagul. Fibrinolysis 4:273-280(1993).
RN [23]
RP VARIANT PROC DEFICIENCY TRP-57.
RX MEDLINE=93271396; PubMed=8499568;


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FT DISULFID 146 159 BY SIMILARITY.
FT DISULFID 161 174 BY SIMILARITY.
FT DISULFID 182 320 INTERCHAIN (BY SIMILARITY).
FT DISULFID 239 255 BY SIMILARITY.
FT DISULFID 373 387 BY SIMILARITY.
FT DISULFID 398 426 BY SIMILARITY.
FT CARBOHYD 215 215 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 291 291 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 355 355 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 461 AA; 51912 MW; 8A4CF93664EDACD5 CRC64;

Query Match 70.6%; Score 139; DB 1; Length 461;
Best Local Similarity 59.1%; Pred. No. 1e-15;
Matches 26; Conservative 7; Mismatches 11; Indels 0; Gaps 0;

QY 1 ANSFLXXLRGSLKXCIXXICDFAKXKIFEDVDVDTLAFWSKH 44
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DB 42 ANSFLFEVRAGSLERSCMEICDFEFAQSLFQNVEDTLAFWKY 85

RESULT 4
ID PRTC RABIT STANDARD; PRT; 458 AA.
AC Q28661;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DE Vitamin-K-dependent protein C precursor (EC 3.4.21.69)
DE (Autoprothrombin IIA) (Anticoagulant protein C) (Blood coagulation factor XIV) (Fragment).
DE PROC.
GN Oryctolagus cuniculus (Rabbit).
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxId=9986;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Shen L., He X., Dahlback B.;
RL Submitted (FEB-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Protein C is a vitamin K-dependent serine protease that
CC regulates blood coagulation by inactivating factors Va and VIIIa
CC in the presence of calcium ions and phospholipids.
CC -!- CATALYTIC ACTIVITY: Degradation of blood coagulation factors Va
CC and VIIIa.
CC -!- SUBUNIT: Synthesized as a single chain precursor, which is cleaved
CC into a light chain and a heavy chain held together by a disulfide
CC bond. The enzyme is then activated by thrombin, which cleaves a
CC tetradecapeptide from the amino end of the heavy chain; this
CC reaction, which occurs at the surface of endothelial cells, is
CC strongly promoted by thrombomodulin.
CC -!- TISSUE SPECIFICITY: Plasma; synthesized in the liver.
CC -!- PTM: The vitamin K-dependent, enzymatic carboxylation of some Glu
CC residues allows the modified protein to bind calcium.
CC -!- MISCELLANEOUS: Calcium also binds, with stronger affinity to
CC another site, beyond the GLA domain. This GLA-independent binding
CC site is necessary for the recognition of the thrombin-
CC thrombomodulin complex.
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 EGF-like domains.
CC -----
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CC -----
CC EMBL; U49933; AAA92956.1;
CC HSSP; P04070; 1PCU.
CC MEROPS; S01.218;
CC InterPro; IPR000152; Asx_hydroxyl_S.

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DR InterPro; IPR009003; Cys_Ser_trypsin.
DR InterPro; IPR001881; EGF_Ca.
DR InterPro; IPR006209; EGF like.
DR InterPro; IPR002383; GLA blood.
DR InterPro; IPR006210; IEGF.
DR InterPro; IPR001254; Peptidase S1.
DR InterPro; IPR001314; Peptidase_SIA.
DR InterPro; IPR000294; VitK_dep_GLA.
DR Pfam; PF00008; EGF; 2.
DR Pfam; PF00594; gla; 1.
DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; SM00001; GLABLOOD.
DR SMART; SM00181; EGF; 2.
DR SMART; SM00089; GLA; 1.
DR SMART; SM00020; Tryp_Spc; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS00026; EGF_3; 1.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE; PS0240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
KW Blood coagulation; Glycoprotein; Serine protease;
KW Gamma-carboxyglutamic acid; Calcium-binding; Vitamin K; Hydroxylation;
FT NON_TER 1
FT SIGNAL <1 27 BY SIMILARITY.
FT PROPEP 28 36 BY SIMILARITY.
FT CHAIN 37 458 VITAMIN K-DEPENDENT PROTEIN C.
FT CHAIN 37 192 PROTEIN C LIGHT CHAIN (BY SIMILARITY).
FT CHAIN 195 458 PROTEIN C HEAVY CHAIN (BY SIMILARITY).
FT PEPTIDE 195 209 ACTIVATION PEPTIDE (BY SIMILARITY).
FT SITE 209 210 CLEAVAGE (BY THROMBIN) (BY SIMILARITY).
FT DOMAIN 91 126 EGF-LIKE 1.
FT DOMAIN 130 170 SERINE PROTEASE.
FT DOMAIN 210 458 EGF-LIKE 2.
FT MOD_RES 42 42 GAMMA-CARBOXYGLUTAMIC ACID
FT MOD_RES 43 43 (BY SIMILARITY).
FT MOD_RES 50 50 GAMMA-CARBOXYGLUTAMIC ACID
FT MOD_RES 52 52 (BY SIMILARITY).
FT MOD_RES 55 55 GAMMA-CARBOXYGLUTAMIC ACID
FT MOD_RES 56 56 (BY SIMILARITY).
FT MOD_RES 61 61 GAMMA-CARBOXYGLUTAMIC ACID
FT MOD_RES 62 62 (BY SIMILARITY).
FT MOD_RES 65 65 GAMMA-CARBOXYGLUTAMIC ACID
FT MOD_RES 107 107 HYDROXYLATION (BY SIMILARITY).
FT ACT_SITE 250 250 CHARGE RELAY SYSTEM.
FT ACT_SITE 296 296 CHARGE RELAY SYSTEM.
FT ACT_SITE 399 399 CHARGE RELAY SYSTEM.
FT DISULFID 53 58 BY SIMILARITY.
FT DISULFID 86 105 BY SIMILARITY.
FT DISULFID 95 100 BY SIMILARITY.
FT DISULFID 99 114 BY SIMILARITY.
FT DISULFID 116 125 BY SIMILARITY.
FT DISULFID 134 145 BY SIMILARITY.
FT DISULFID 141 154 BY SIMILARITY.
FT DISULFID 156 169 BY SIMILARITY.
FT DISULFID 177 316 INTERCHAIN (BY SIMILARITY).
FT DISULFID 235 251 BY SIMILARITY.
FT DISULFID 370 384 BY SIMILARITY.
FT DISULFID 395 423 BY SIMILARITY.

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FT DISULFID 396 424 BY SIMILARITY.
FT CARBOHYD 138 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 292 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 353 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 459 AA; 51866 MW; 8541AAC14CCL6D09 CRC64;

Query Match 62.4%; Score 123; DB 1; Length 459;
Best Local Similarity 52.3%; Pred. No. 5.4e-13;
Matches 23; Conservative 7; Mismatches 14; Indels 0; Gaps 0;

QY 1 ANSFLXLRGSLXKXICXIXICDFXAKXIFEDVDDTLAWSH 44
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 42 ANSFLLELPSSLERECKTCTDFEAREIFQNTMTAFWSKY 85

RESULT 6
PRTC_BOVIN STANDARD; PRT; 456 AA.
AC P007A5;
DT 21-JUL-1986 (Rel. 01, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Vitamin-K-dependent protein C precursor (PC 3, 4, 21-69)
DE (Autoproteolysis IIA) (Anticoagulant protein C) (Blood coagulation
DE factor XIV) (Fragment).
DE PROC.
GN Bos taurus (Bovine).
OC Bos taurus; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=85014826; PubMed=6091100;
RA Long G.L., Balagaje R.M., McGillivray R.T.A.;
RT "Cloning and sequencing of liver cDNA coding for bovine protein C.";
RL Proc. Natl. Acad. Sci. U.S.A. 81:5653-5656(1984).
RN [2]
RP SEQUENCE OF 40-194, AND CARBOHYDRATE-LINKAGE SITE ASN-136.
RX MEDLINE=83007325; PubMed=6896876;
RA Fernlund P., Stenflo J.;
RT "Amino acid sequence of the light chain of bovine protein C.";
RL J. Biol. Chem. 257:12170-12179(1982).
RN [3]
RP REVISION TO 110.
RX MEDLINE=83169769; PubMed=6572939;
RA Drakenberg T., Fernlund P., Roepstorff P., Stenflo J.;
RT "Beta-hydroxyaspartic acid in vitamin K-dependent protein C.";
RL Proc. Natl. Acad. Sci. U.S.A. 80:1802-1806(1983).
RN [4]
RP SEQUENCE OF 197-456, AND CARBOHYDRATE-LINKAGE SITES ASN-289; ASN-350
RX AND ASN-366.
RX MEDLINE=83007326; PubMed=6896877;
RA Stenflo J., Fernlund P.;
RT "Amino acid sequence of the heavy chain of bovine protein C.";
RL J. Biol. Chem. 257:12180-12190(1982).
RN [5]
RP PROCESSING, AND CALCIUM-BINDING DATA.
RX MEDLINE=83213513; PubMed=6304032;
RA Esmon N.L., Debault L.S., Esmon C.T.;
RT "Proteolytic formation and properties of gamma-carboxyglutamic acid-
RT domainless protein C.";
RL J. Biol. Chem. 258:5548-5553(1983).
RN [6]
RP PROCESSING, AND CALCIUM-BINDING DATA.
RX MEDLINE=83213514; PubMed=6406503;
RA Johnson A.E., Esmon N.L., Laue T.M., Esmon C.T.;
RT "Structural changes required for activation of protein C are induced
RT by Ca2+ binding to a high affinity site that does not contain gamma-
RT carboxyglutamic acid.";
RL J. Biol. Chem. 258:5554-5560(1983).
CC -!- FUNCTION: Protein C is a vitamin K-dependent serine protease that
    regulates blood coagulation by inactivating factors Va and VIIIa

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CC in the presence of calcium ions and phospholipids.
CC -!- CATALYTIC ACTIVITY: Degradation of blood coagulation factors Va
CC and VIIIa.
CC -!- SUBUNIT: Synthesized as a single chain precursor, which is cleaved
CC bond. The enzyme is then activated by thrombin, which cleaves a
CC tetradecapeptide from the amino end of the heavy chain; this
CC reaction, which occurs at the surface of endothelial cells, is
CC strongly promoted by thrombomodulin.
CC -!- TISSUE SPECIFICITY: Plasma; synthesized in the liver.
CC -!- PTM: The vitamin K-dependent, enzymatic carboxylation of some Glu
CC residues allows the modified protein to bind calcium.
CC -!- MISCELLANEOUS: Calcium also binds, with stronger affinity to
CC another site, beyond the GLA domain. This GLA-independent binding
CC site is necessary for the recognition of the thrombin-
CC thrombomodulin complex.
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 EGF-like domains.
CC -----
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CC -----
CC EMBL; K02435; AAA30685.1; -.
CC PIR; A26250; KXBO.
CC HSRP; P04070; LPCU.
CC MEROPS; S01.218; -.
CC InterPro; IPR000152; Asx_Hydroxyl_S.
CC InterPro; IPR009003; Cys_Ser_trypsin.
CC InterPro; IPR001881; EGF_Ca.
CC InterPro; IPR006209; EGF_Like.
CC InterPro; IPR002383; GLA_Blood.
CC InterPro; IPR006210; IEGF.
CC InterPro; IPR001254; Peptidase_S1.
CC InterPro; IPR001314; Peptidase_S1A.
CC InterPro; IPR000294; VitK_dep_GLA.
CC Pfam; PF00008; EGF_2.
CC Pfam; PF00594; Gla_1.
CC Pfam; PF00083; tryptain; 1.
CC PRINTS; PR00722; CHYMOTRYPSIN.
CC PRINTS; PR00001; GLABLOOD.
CC SMART; SM00181; EGF_2.
CC SMART; SM00069; GLA; 1.
CC SMART; SM00020; Tryp_Spc; 1.
CC PROSITE; PS00010; ASX_HYDROXYL; 1.
CC PROSITE; PS00022; EGF_1; 1.
CC PROSITE; PS01186; EGF_2; 2.
CC PROSITE; PS00026; EGF_3; 1.
CC PROSITE; PS01187; EGF_CA; 1.
CC PROSITE; PS00011; GLU_CARBOXYLATION; 1.
CC PROSITE; PS02040; TRYPSIN_DOM; 1.
CC PROSITE; PS00134; TRYPSIN_HIS; FALSE_NEG.
CC PROSITE; PS00135; TRYPSIN_SER; 1.
CC Blood coagulation; Glycoprotein; Serine protease;
CC Gamma-carboxyglutamic acid; Calcium-binding; Vitamin K; Hydroxylation;
CC EGF-like domain; Repeat; Endothelial cell; Hydrolase; Signal.
CC NON_TER 1 1
CC SIGNAL <1 29
CC PROPEP 30 39
CC CHAIN 40 194
CC CHAIN 197 456
CC PEPTIDE 197 210
CC DOMAIN 94 129
CC DOMAIN 133 173
CC DOMAIN 211 456
CC MOD_RES 45 45
CC MOD_RES 46 46
CC MOD_RES 53 53
CC MOD_RES 55 55
CC PROTEIN C LIGHT CHAIN.
CC PROTEIN C HEAVY CHAIN.
CC ACTIVATION PEPTIDE.
CC EGF-Like 1.
CC EGF-Like 2.
CC SERINE PROTEASE.
CC GAMMA-CARBOXYGLUTAMIC ACID.
CC GAMMA-CARBOXYGLUTAMIC ACID.
CC GAMMA-CARBOXYGLUTAMIC ACID.

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FT MOD RES 58 GAMMA-CARBOXYGLUTAMIC ACID.
FT MOD RES 59 GAMMA-CARBOXYGLUTAMIC ACID.
FT MOD RES 62 GAMMA-CARBOXYGLUTAMIC ACID.
FT MOD RES 64 GAMMA-CARBOXYGLUTAMIC ACID.
FT MOD RES 65 GAMMA-CARBOXYGLUTAMIC ACID.
FT MOD RES 68 GAMMA-CARBOXYGLUTAMIC ACID.
FT MOD RES 74 GAMMA-CARBOXYGLUTAMIC ACID.
FT MOD RES 110 HYDROXYLATION.
FT ACT SITE 252 CHARGE RELAY SYSTEM.
FT ACT SITE 298 CHARGE RELAY SYSTEM.
FT ACT SITE 397 CHARGE RELAY SYSTEM.
FT DISULFID 56 BY SIMILARITY.
FT DISULFID 89 BY SIMILARITY.
FT DISULFID 98 BY SIMILARITY.
FT DISULFID 102 BY SIMILARITY.
FT DISULFID 119 BY SIMILARITY.
FT DISULFID 137 BY SIMILARITY.
FT DISULFID 144 BY SIMILARITY.
FT DISULFID 159 BY SIMILARITY.
FT DISULFID 180 INTERCHAIN.
FT DISULFID 237
FT DISULFID 368
FT DISULFID 393
FT CARBOHYD 136 N-LINKED (GLCNAC. . .)
FT CARBOHYD 289 N-LINKED (GLCNAC. . .)
FT CARBOHYD 350 N-LINKED (GLCNAC. . .)
FT CARBOHYD 366 F -> K.
FT VARIANT 82 VP -> PV (IN REF. 4).
FT CONFLICT 455
SQ SEQUENCE 456 AA; 51407 MW; CAAFG833F894C209 CRC64;

Query Match 61.9%; Score 122; DB 1; Length 456;
Best Local Similarity 50.0%; Pred. No. 7.9e-13;
Matches 21; Conservative 9; Mismatches 12; Indels 0; Gaps 0;

QY 1 ANSFLXLRQSLRXRCITXICDFXXAKXIFEDVDLTFWS 42
DB 40 ANSFLRLPGNVERCSEECVEFEAREIFQNTEDTAFWS 81

RESULT 7
FA10_BOVIN STANDARD; PRT; 492 AA.
AC P00743;
DT 21-JUL-1986 (Rel. 01, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Coagulation factor X precursor (EC 3.4.21.6) (Stuart factor).
GN F10.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE OF 1-487 FROM N.A.
RX MEDLINE=84247315; PubMed=6330671;
RA Fung M.R., Campbell R.M., McGillicray R.T.A.;
RT "Blood coagulation factor X mRNA encodes a single polypeptide chain
containing a propro leader sequence."
RL Nucleic Acids Res. 12:4481-4492(1984).
RN [2]
RP SEQUENCE OF 41-180.
RX MEDLINE=80130563; PubMed=6766735;
RA Enfield D.L., Ericsson L.H., Fujikawa K., Walsh K.A., Neurath H.,
RA Titani K.;
RT "Amino acid sequence of the light chain of bovine factor X1 (Stuart
factor).";
RL Biochemistry 19:659-667(1980).
RN [3]
RP REVISION TO 103.
RX MEDLINE=83308813; PubMed=6688526;
RA McMullen B.A., Fujikawa K., Kistiel W.;

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RT "The occurrence of beta-hydroxyaspartic acid in the vitamin
K-dependent blood coagulation zymogens."
RL Biochem. Biophys. Res. Commun. 115:8-14(1983).
RN [4]
RP SEQUENCE OF 183-492, CARBOHYDRATE-LINKAGE SITES, AND DISULFIDE BONDS.
RX MEDLINE=760533069; PubMed=1059093;
RA Titani K., Fujikawa K., Enfield D.L., Ericsson L.H., Walsh K.A.,
RA Neurath H.;
RT "Bovine factor X1 (Stuart factor): amino-acid sequence of heavy
chain.";
RL Proc. Natl. Acad. Sci. U.S.A. 72:3082-3086(1975).
RN [5]
RP SEQUENCE OF 183-233, AND CARBOHYDRATE-LINKAGE SITES.
RX MEDLINE=94062825; PubMed=8243451;
RA Inoue K., Morita T.;
RT "Identification of O-linked oligosaccharide chains in the activation
peptides of blood coagulation factor X. The role of the carbohydrate
moieties in the activation of factor X.";
RL Eur. J. Biochem. 218:153-163(1993).
RN [6]
RP ACTIVE SITE.
RX MEDLINE=73053314; PubMed=4364286;
RA Titani K., Hermodeen M.A., Fujikawa K., Ericsson L.H., Walsh K.A.,
RA Neurath H., Davie E.W.;
RT "Bovine factor X1a (activated Stuart factor). Evidence of homology
with mammalian serine proteases.";
RL Biochemistry 11:4899-4903(1972).
RN [7]
RP PROCESSING.
RX MEDLINE=76053121; PubMed=1059122;
RA Fujikawa K., Titani K., Davie E.W.;
RT "Activation of bovine factor X (Stuart factor): conversion of factor
Xa-alpha to factor Xa-beta.";
RL Proc. Natl. Acad. Sci. U.S.A. 72:3359-3363(1975).
RN [8]
RP CALCIUM-BINDING DATA.
RX MEDLINE=84185716; PubMed=6546930;
RA Sugo I., Björk I., Holmgren A., Stenflo J.;
RT "Calcium-binding properties of bovine factor X lacking the gamma-
carboxyglutamic acid-containing region.";
RL J. Biol. Chem. 259:5705-5710(1984).
RN [9]
RP SULFATION.
RX MEDLINE=86140210; PubMed=3949800;
RA Morita T., Jackson C.M.;
RT "Localization of the structural difference between bovine blood
coagulation factors X1 and X2 to tyrosine 18 in the activation
peptide.";
RL J. Biol. Chem. 261:4008-4014(1986).
RN [10]
RP STRUCTURE BY NMR OF 85-126.
RX MEDLINE=91084483; PubMed=2261466;
RA Selander M., Persson E., Stenflo J., Drakenberg T.;
RT "1H NMR assignment and secondary structure of the Ca2(+)-free form of
the amino-terminal epidermal growth factor like domain in coagulation
factor X.";
RL Biochemistry 29:8111-8118(1990).
RN [11]
RP STRUCTURE BY NMR OF 85-126.
RX MEDLINE=92329412; PubMed=1627540;
RA Ullner M., Selander M., Persson E., Stenflo J., Drakenberg T.,
RA Teleman O.;
RT "Three-dimensional structure of the apo form of the N-terminal
EGF-like module of blood coagulation factor X as determined by NMR
spectroscopy and simulated folding.";
RL Biochemistry 31:5974-5983(1992).
RN [12]
RP STRUCTURE BY NMR OF 85-126.
RX MEDLINE=92406922; PubMed=1527084;
RA Selander-Sunnerhagen M., Ullner M., Persson E., Teleman O.,
RA Stenflo J., Drakenberg T.;
RT "How an epidermal growth factor (EGF)-like domain binds calcium. High
resolution NMR structure of the calcium form of the NH2-terminal EGF-

```

RT like domain in coagulation factor X.";

RL J. Biol. Chem. 267:19642-19649(1992).

RN [13]

RP STRUCTURE BY NMR OF 41-126.

RX MEDLINE=96387194; PubMed=8794734;

RA Sunnerhagen M., Olah G.A., Stenflo J., Forsen S., Drakenberg T.,

RA Trewhella J.;

RT "The relative orientation of Gla and EGF domains in coagulation

RT factor X is altered by Ca²⁺ binding to the first EGF domain. A

RT combined NMR-small angle X-ray scattering study.";

RL Biochemistry 35:11547-11559(1996).

CC CC -1- FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that

CC converts prothrombin to thrombin in the presence of factor Va,

CC calcium and phospholipid during blood clotting.

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CC Arg-|-Ile bonds in prothrombin to form thrombin.

CC CC -1- SUBUNIT: The two chains are formed from a single-chain precursor

CC by the exclusion of two Arg residues and are held together by 1 or

CC more disulfide bonds.

CC CC -1- PTM: The vitamin K-dependent, enzymatic carboxylation of some

CC glutamate residues allows the modified protein to bind calcium.

CC CC -1- PTM: N- and O-glycosylated.

CC CC -1- PTM: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE

CC INTRINSIC PATHWAY), OR BY FACTOR VIIA (IN THE EXTRINSIC PATHWAY).

CC CC -1- MISCELLANEOUS: Calcium also binds, with stronger affinity to

CC another site, beyond the GLA domain.

CC CC -1- SIMILARITY: Belongs to peptidase family S1.

CC CC -1- SIMILARITY: Contains 2 EGF-like domains.

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CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>

CC or send an email to license@isb-sib.ch).

CC -----

DR EMBL; X00673; CA25286.1; -;

DR FIR; A22867; EXBO.

DR PDB; 1AFO; 31-JAN-94.

DR PDB; 1CCF; 31-MAY-94.

DR PDB; 1WHE; 15-MAY-97.

DR PDB; 1WHF; 15-MAY-97.

DR PDB; 1IOD; 21-JAN-03.

DR PDB; 1KIG; 28-OCT-98.

DR MEROPS; S01.216; -;

DR GlycoSuiteDB; P00743; -;

DR InterPro; IPR000152; Asx_hydroxyl_S.

DR InterPro; IPR009003; Cys_ser_trypsin.

DR InterPro; IPR000742; EGF_2.

DR InterPro; IPR001881; EGF_Ca.

DR InterPro; IPR006209; EGF_like.

DR InterPro; IPR002383; GLA_blood.

DR InterPro; IPR001254; Peptidase_S1.

DR InterPro; IPR001314; Peptidase_S1A.

DR InterPro; IPR000294; VitK_dep_GLA.

DR Pfam; PF00008; EGF; 2.

DR Pfam; PF00594; Gla; 1.

DR Pfam; PF00089; trypsin; 1.

DR PRINTS; PR00722; CHYNOTRYPsin.

DR PRINTS; PR00001; GLABLOOD.

DR SMART; SM00179; EGF_CA; 1.

DR SMART; SM00069; GLA; 1.

DR SMART; SM00020; TRYD_SPG; 1.

DR PROSITE; PS00010; ASX_HYDROXYL; 1.

DR PROSITE; PS00022; EGF_1; 1.

DR PROSITE; PS01186; EGF_2; 2.

DR PROSITE; PS00026; EGF_3; 1.

DR PROSITE; PS01187; EGF_CA; 1.

DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.

DR PROSITE; PS00240; TRYPSIN_DOM; 1.

DR PROSITE; PS00134; TRYPSIN_HIS; 1.

DR PROSITE; PS00135; TRYPSIN_SER; 1.

KW Glycoprotein; Hydrolase; Serine protease; Plasma; Blood coagulation;

KW Gamma-carboxyglutamic acid; Hydroxylation; Calcium-binding; Vitamin K;

KW Signal; Zymogen; EGF-like domain; Repeat; Sulfation; 3D-structure.

FT SIGNAL 1 23 POTENTIAL.

FT PROPEP 24 40 FACTOR X LIGHT CHAIN.

FT CHAIN 41 180 FACTOR X HEAVY CHAIN.

FT CHAIN 183 492 ACTIVATION PEPTIDE.

FT PROPEP 183 233 ACTIVATED FACTOR XA, HEAVY CHAIN.

FT CHAIN 234 492 MAY BE REMOVED BUT IS NOT NECESSARY FOR

FT PROPEP 476 492 ACTIVATION.

FT DOMAIN 86 122 EGF-LIKE 1, CALCIUM-BINDING (POTENTIAL).

FT DOMAIN 125 165 EGF-LIKE 2,

FT DOMAIN 234 492 SERINE PROTEASE.

FT ACT_SITE 275 275 CHARGE RELAY SYSTEM.

FT ACT_SITE 321 321 CHARGE RELAY SYSTEM.

FT ACT_SITE 418 418 CHARGE RELAY SYSTEM.

FT MOD_RES 46 46 GAMMA-CARBOXYGLUTAMIC ACID.

FT MOD_RES 47 47 GAMMA-CARBOXYGLUTAMIC ACID.

FT MOD_RES 54 54 GAMMA-CARBOXYGLUTAMIC ACID.

FT MOD_RES 56 56 GAMMA-CARBOXYGLUTAMIC ACID.

FT MOD_RES 59 59 GAMMA-CARBOXYGLUTAMIC ACID.

FT MOD_RES 60 60 GAMMA-CARBOXYGLUTAMIC ACID.

Query Match 57.9%; Score 114; DB 1; Length 492;

Best Local Similarity 45.5%; Pred. No. 2e-11; Mismatches 16; Indels 0; Gaps 0;

Matches 20; Conservative 8; Mismatches 16; Indels 0; Gaps 0;

QY 1 ANSFLLXLRQSLNRXIXICDFXAKXIFEDVDVDTLAFWSKH 44

DB 41 ANSFLEBVKQNLRECLERACSLERAEVFEAEQTDFFWSKY 84

RESULT 8

FA10_HUMAN

ID FA10_HUMAN STANDARD; PRT; 488 AA.

AC P00742; Q14340;

DT 21-JUL-1986 (Rel. 01, Created)

DT 01-OCT-1989 (Rel. 12, Last sequence update)

DT 10-OCT-2003 (Rel. 42, Last annotation update)

DE Coagulation factor X precursor (EC 3.4.21.6) (Stuart factor).

GN F10.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=91216473; PubMed=1902434;

RA Messier T.L., Pittman D.D., Long G.L., Kaufman R.J., Church W.R.;

RT "Cloning and expression in COS-1 cells of a full-length cDNA encoding

human coagulation factor X.";

RL Gene 99:291-294(1991).

RN [2]

RP SEQUENCE FROM N.A.

RX MEDLINE=87026600; PubMed=3768336;

RA Leytus S.P., Foster D.C., Kurachi K., Davie E.W.;

RT "Gene for human factor X: a blood coagulation factor whose gene

organization is essentially identical with that of factor IX and

protein C.";

RL Biochemistry 25:5098-5102(1986).

RN [3]

RP SEQUENCE FROM N.A.

RX MEDLINE=22388257; PubMed=12477932;

RA Strausberg R.L., Feigold E.A., Grouse L.H., Derge J.G.,

RA Klausner R.D., Collins F.S., Wagner L., Shennan C.M., Schuler G.D.,

RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,

RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Narusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.D., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carinacci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Woxley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Kryzysinski M.I., Skalska U., Smallus D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.,
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences".
EL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [5]
RN SEQUENCE OF 13-488 FROM N.A.
RN MEDLINE=85216545; PubMed=2582420;
RA Fung M.R., Hay C.W., McGillivray R.T.A.,
RT "Characterization of an almost full-length cDNA coding for human
RT blood coagulation factor X".
EL Proc. Natl. Acad. Sci. U.S.A. 82:3591-3595(1985).
RN [6]
RN SEQUENCE OF 19-488 FROM N.A.
RN TISSUE=Liver;
RN MEDLINE=86221713; PubMed=3011603;
RA Kaul R.K., Hildebrand B., Roberts S., Jagadeeswaran P.,
RT "Isolation and characterization of human blood-coagulation factor X
RT cDNA".
EL Gene 41:311-314(1986).
RN [7]
RN SEQUENCE OF 41-179.
RN MEDLINE=93257207; PubMed=6871167;
RA McMullen B.A., Fujikawa K., Kistiel W., Sasagawa T., Howald W.N.,
RA Kwa E.Y., Weinstein B.,
RT "Complete amino acid sequence of the light chain of human blood
RT coagulation factor X: evidence for identification of residue 63 as
RT beta-hydroxyaspartic acid".
EL Biochemistry 22:2875-2884(1983).
RN [8]
RN SEQUENCE OF 115-488 FROM N.A., AND TISSUE SPECIFICITY.
RN TISSUE=Liver;
RN MEDLINE=94062825; PubMed=8243461;
RA Inoue K., Morita T.,
RT "Identification of O-linked oligosaccharide chains in the activation
RT peptides of blood coagulation factor X. The role of the carbohydrate
RT moieties in the activation of factor X".
EL Eur. J. Biochem. 218:153-163(1993).
RN [10]
RN SEQUENCE OF 1-23 FROM N.A.
RN MEDLINE=90128299; PubMed=2612918;
RA Jagadeeswaran P., Reddy S.V., Rao K.J., Hamsabhusanam K., Lyman G.,
RT "Cloning and characterization of the 5' end (exon 1) of the gene
RT encoding human factor X".
EL Gene 84:517-519(1989).
RN [11]
RN X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 86-179 AND 235-278.
RN MEDLINE=93360277; PubMed=8355279;
RA Padmanabhan K., Padmanabhan K.P., Tulinsky A., Park C.H., Bode W.,
RA Huber R., Blankenship D.T., Cardin A.D., Kistiel W.,
RT "Structure of human des(1-45) factor Xa at 2.2-A resolution".
EL J. Mol. Biol. 232:947-966(1993).
RN [12]
RN X-RAY CRYSTALLOGRAPHY (2.3 ANGSTROMS) OF 86-179 AND 235-278.
RN MEDLINE=98283982; PubMed=9618463;
RA Kamata K., Kawamoto H., Honma T., Iwama T., Kim S.H.,
RT "Structural basis for chemical inhibition of human blood coagulation
RT factor Xa".
EL Proc. Natl. Acad. Sci. U.S.A. 95:6630-6635(1998).
RN [13]
RN VARIANTS ILE-7 AND HIS-30.
RN MEDLINE=99318093; PubMed=10391209;
RA Cargill M., Altshuler D., Ireland J., Sklar P., Ardlie K., Patil N.,
RA Shaw N., Lane C.R., Lim E.P., Kalyanaram N., Nemesh J., Ziaugra L.,
RA Friedland L., Rolfe A., Warrington J., Lipshutz R., Daley G.Q.,
RA Lander E.S.,
RT "Characterization of single-nucleotide polymorphisms in coding regions
RT of human genes".
EL Nat. Genet. 22:231-238(1999).
RN [14]
RN ERRATUM.
RA Cargill M., Altshuler D., Ireland J., Sklar P., Ardlie K., Patil N.,
RA Shaw N., Lane C.R., Lim E.P., Kalyanaram N., Nemesh J., Ziaugra L.,
RA Friedland L., Rolfe A., Warrington J., Lipshutz R., Daley G.Q.,
RA Lander E.S.,
RL Nat. Genet. 23:373-373(1999).
CC -!- FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that
CC converts prothrombin to thrombin in the presence of factor Va,
CC calcium and phospholipid during blood clotting.
CC -!- CATALYTIC ACTIVITY: Preferential cleavage: Arg-|-Thr and then
CC Arg-|-Ile bonds in prothrombin to form thrombin.
CC -!- SUBUNIT: The two chains are formed from a single-chain precursor
CC by the excision of two Arg residues and are held together by 1 or
CC more disulfide bonds.
CC -!- TISSUE SPECIFICITY: Plasma; synthesized in the liver.
CC -!- PTM: The vitamin K-dependent, enzymatic carboxylation of some
CC glutamate residues allows the modified protein to bind calcium.
CC -!- PTM: N- and O-glycosylated.
CC -!- PTM: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE
CC INTRINSIC PATHWAY), OR BY FACTOR VIIA (IN THE EXTRINSIC PATHWAY).
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 EGF-like domains.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
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CC entities requires a license agreement (see [http://www.isb-sib.ch/](http://www.isb-sib.ch/announce/)
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; K03194; AAA52490.1; -
CC EMBL; M57285; AAA52421.1; -
CC EMBL; AF503510; AM19347.1; -
CC EMBL; BC046125; AAH46125.1; -
CC EMBL; L29433; AAA52764.1; -
CC EMBL; L00390; AAA52764.1; JOINED.
CC EMBL; L00391; AAA52764.1; JOINED.
CC EMBL; L00392; AAA52764.1; JOINED.
CC EMBL; L00393; AAA52764.1; JOINED.
CC EMBL; L00394; AAA52764.1; JOINED.
CC EMBL; L00395; AAA52764.1; JOINED.
CC EMBL; L00396; AAA52764.1; JOINED.
CC EMBL; M22613; AAA51984.1; -
CC EMBL; K01886; AAA52486.1; -
CC EMBL; M33297; AAA52636.1; -
CC PIR; A24478; EXHU.
CC PDB; 1HCG; 08-MAY-95.
CC PDB; 1FAF; 23-OCT-97.
CC PDB; 1FXV; 17-JUN-98.
CC PDB; 1XKA; 23-MAR-99.
CC PDB; 1XKB; 23-MAR-99.
CC PDB; 1EZO; 20-SEP-00.
CC PDB; 1F08; 20-SEP-00.
CC PDB; 1F08; 20-SEP-00.
CC PDB; 1F0S; 17-NOV-00.
CC PDB; 1G2L; 20-OCT-01.
CC PDB; 1G2M; 20-OCT-01.

PROTEIN 3.

EXTRACELLULAR (POTENTIAL).
POTENTIAL.
CYTOPLASMIC (POTENTIAL).
GLA-RICH.

8A373E48490D61 CRC64;

Score 107; DB 1; Length 231;
Best Local Similarity 43.9%; Pred.No.1.3e-10;
Matches 18; Conservative S; Mismatches 15; Indels 0; Gaps

Oy 1 ANSFLXLRQGSIXKXCXXCICDPFXXAKXFEDVDDTLAFW 41
| | : ~~~~~~
Db 20 ANEFLELROGTTIERECWEIICSVEEVKEVFENKEMTFEW 60
| | : ~~~~~~

RESULT 10

FA10 RABIT STANDARD; PRT; 490 AA.

ID FA10 RABIT

OS Oryctolagus cuniculus (Rabbit).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.

NCBI_TaxID=9986;

[]]

RN SEQUENCE FROM N.A.

RP MEDLINE=97256311; PubMed=9101642;

RX Pandurith U.R., Anderson K.D.; James H.L.;

RA "Characterization of a full-length CDNA for rabbit factor X.";

RL Thromb. Res. 85:503-514(1997)

CC -! FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that converts prothrombin to thrombin in the presence of factor Va, calcium and phospholipid during blood clotting.

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CC -! SUBUNIT: The two chains are formed from a single-chain precursor by the excision of two Arg residues and are held together by 1 or more disulfide bonds.

CC -! PTM: The vitamin K-dependent, enzymatic carboxylation of some glutamate residues allows the modified protein to bind calcium (By similarity).

CC -! PTM: N- and O-glycosylated (By similarity).

CC -! PTM: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE INTRINSIC PATHWAY), OR BY FACTOR VIIIA (IN THE EXTRINSIC PATHWAY) (BY SIMILARITY).

CC -! MISCELLANEOUS: Calcium also binds, with stronger affinity to another site, beyond the GLA domain.

CC -! SIMILARITY: Belongs to peptidase family S1.

CC -! SIMILARITY: Contains 2 EGF-like domains.

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EMBL; AF003200; AAB62542.1; -.
HSSP; F00742; LHCG.
MEROPS; S01.216; .
InterPro; IPR000152; Asx_hydroxyl_S.
InterPro; IPR009003; Cys_Ser_trypsin.
InterPro; IPR000742; EGF_2.
InterPro; IPR001881; EGF_F2.
InterPro; IPR001438; EGF II.
InterPro; IPR006209; EGF like.
InterPro; IPR002383; GLA_blood.

DR InterPro; IPR001254; Peptidase S1.
 DR InterPro; IPR001314; Peptidase-S1A.
 DR InterPro; IPR000294; Vtck_dep_GLA.
 DR Pfam; PF00008; EGF; 2.
 DR Pfam; PF00594; Gla; 1.
 DR PRINTS; PRO0722; trypsin; 1.
 DR PRINTS; PRO0722; CHYMOTRYPSIN.
 DR PRINTS; PRO0010; EGFBLD.
 DR PRINTS; PRO0001; GLABLOOD.
 DR SMART; SM00179; EGF_CA; 1.
 DR SMART; SM00069; GLA; 1.
 DR SMART; SM00020; Tryp_Spc; 1.
 DR PROSITE; PS00010; ASX_HYDROXYL; 1.
 DR PROSITE; PS00022; EGF_1; 1.
 DR PROSITE; PS01186; EGF_2; 2.
 DR PROSITE; PS00026; EGF_3; 1.
 DR PROSITE; PS01187; EGF_CA; 1.
 DR PROSITE; PS00011; GLU_CARBOXYLATION; 1.
 DR PROSITE; PS00240; TRYPSIN_DOM; 1.
 DR PROSITE; PS00134; TRYPSIN_HIS; 1.
 DR PROSITE; PS00135; TRYPSIN_SER; 1.
 KW Glycoprotein; Hydrolase; Serine protease; Plasma; Blood coagulation;
 KW Gamma-carboxyglutamic acid; Hydroxylation; Calcium-binding; Vitamin K;
 KW Signal; Zymogen; EGF-like domain; Repeat.
 FT SIGNAL 1 20
 FT PROPEP 21 40
 FT CHAIN 41 180
 FT CHAIN 184 490
 FT PROPEP 184 232
 FT CHAIN 233 490
 FT DOMAIN 86 122
 FT DOMAIN 125 165
 FT DOMAIN 233 490
 FT MOD_RES 46 46
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 FT MOD_RES 103 103
 FT ACT_SITE 274 274
 FT ACT_SITE 320 320
 FT ACT_SITE 417 417
 FT DISULFID 90 101
 FT DISULFID 95 110
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 FT DISULFID 151 164
 FT DISULFID 172 340
 FT DISULFID 239 244
 FT DISULFID 259 275
 FT DISULFID 388 402
 FT DISULFID 413 441
 FT CARBOHYD 61 61
 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 187 187 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 205 205 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 490 AA; 53965 MW; 3A39FA85AF2A6D11 CRC64;
 Query Match 52.3%; Score 103; DB 1; Length 490;
 Best Local Similarity 43.2%; Pred No. 1.4e-09;
 Matches 19; Conservative 9; Mismatches 16; Indels 0; Gaps 0;
 QY 1 ANSFLXLRQGSLLXRCXICDFXXAXKXIFEDVDTDLAFWSKH 44
 DB 41 ANSFLEELKGNLERECMEENGCSYEALVEFEDREKTNFVWKY 84
 RESULT 11
 PA7_RABIT STANDARD; PRT; 444 AA.
 ID AC P98139; P79224;
 DT 01-FEB-1996 (Rel. 33, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 10-OCT-2003 (Rel. 42, Last annotation update)
 DE Coagulation factor VII precursor (EC 3.4.21.21) (Serum prothrombin
 DE conversion accelerator).
 GN F7.
 OS Oryctolagus cuniculus (Rabbit).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
 OX NCBI_TaxID=9986;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RX MEDLINE=93190306; PubMed=8383365;
 RA Brothers A.B., Clarke B.J., Sheffield W.P., Blajchman M.A.;
 RT "Complete nucleotide sequence of the cDNA encoding rabbit coagulation
 RT factor VII.";
 RL Thromb. Res. Suppl. 69:231-238(1993).
 RN [2]
 RP REVISION TO 395.
 RC TISSUE=Liver;
 RA Ruiz S.R., Blajchman M.A., Clarke B.J.;
 RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Circulates in the blood in a zymogen form. Factor VII is
 CC converted to factor VIIa by factor Xa, factor XIa, factor IXa, or
 CC thrombin by minor proteolysis. In the presence of tissue factor
 CC and calcium ions, factor VIIa then converts factor X to factor Xa
 CC by limited proteolysis. Factor VIIa will also convert factor IX to
 CC factor IXa in the presence of tissue factor and calcium (By
 CC similarity).
 CC -!- CATALYTIC ACTIVITY: Hydrolyzes one Arg-|-Ile bond in factor X to
 CC form factor Xa.
 CC -!- SUBUNIT: Heterodimer of a light chain and a heavy chain linked by
 CC a disulfide bond (By similarity).
 CC -!- TISSUE SPECIFICITY: Plasma.
 CC -!- PTM: The vitamin K-dependent, enzymatic carboxylation of some
 CC glutamate residues allows the modified protein to bind calcium (By
 CC similarity).
 CC -!- SIMILARITY: Belongs to peptidase family S1.
 CC -!- SIMILARITY: Contains 2 EGF-like domains.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (see <http://www.isb-sib.ch/announce/>
 CC or send an email to license@sib-sib.ch).
 CC -----
 CC EMBL; U77477; AAB37326.1; -;
 CC HSSP; P08709; 1FAK.
 CC MEROPS; S01.215; -;
 CC InterPro; IPR000152; Asx_hydroxyl_S.
 CC InterPro; IPR009003; Cys_Ser_trypsin.
 CC InterPro; IPR000742; EGF_2.
 CC InterPro; IPR001881; EGF_Ca.
 DR

DR InterPro: IPR001438; EGF II.
DR InterPro: IPR006209; EGF-like.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Peptidase_s1.
DR InterPro: IPR001314; Peptidase_s1A.
DR InterPro: IPR000294; VitK_dep_GLA.
DR Pfam: PF00008; EGF; 2.
DR Pfam: PF00594; gla; 1.
DR Pfam: PF00089; trypsin; 1.
DR PRINTS: PR00722; CHYMOTRYPSIN.
DR PRINTS: PR00010; EGF_blood.
DR PRINTS: PR00001; GLABLOOD.
DR SMART: SM00179; EGF_CA; 1.
DR SMART: SM00069; GLA; 1.
DR SMART: SM00020; Tryp_SPC; 1.
DR PROSITE: PS00010; ASK_HYDROXYL; 1.
DR PROSITE: PS00022; EGF_1; 1.
DR PROSITE: PS01186; EGF_2; 1.
DR PROSITE: PS00026; EGF_3; 1.
DR PROSITE: PS01187; EGF_CA; 1.
DR PROSITE: PS00011; GLU CARBOXYLATION; 1.
DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; 1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
KW Hydrolase; Serine protease; Blood coagulation; Zymogen; Glycoprotein;
KW Liver; Plasma; Vitamin K; Calcium-binding; Gamma-carboxyglutamic acid;
KW EGF-like domain; Repeat; Signal; Hydroxylation.
FT SIGNAL 1 21
FT PROPEP 22 39
FT CHAIN 40 191
FT CHAIN 192 444
FT DOMAIN 45 74
FT DOMAIN 85 121
FT DOMAIN 126 167
FT DOMAIN 192 444
FT SITE 191 192
FT ACT_SITE 232 232
FT ACT_SITE 281 281
FT ACT_SITE 383 383
FT BINDING 377 377
FT DISULFID 56 61
FT DISULFID 89 100
FT DISULFID 111 120
FT DISULFID 130 141
FT DISULFID 137 151
FT DISULFID 153 166
FT DISULFID 174 301
FT DISULFID 198 203
FT DISULFID 217 233
FT DISULFID 349 368
FT DISULFID 379 407
FT MOD_RES 45 45
FT MOD_RES 46 46
FT MOD_RES 53 53
FT MOD_RES 55 55
FT MOD_RES 58 58
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FT MOD_RES 64 64
FT MOD_RES 65 65
FT MOD_RES 68 68
FT MOD_RES 74 74
FT MOD_RES 102 102
FT CARBOHYD 211 211
FT CARBOHYD 242 242
FT CARBOHYD 306 306
SQ SEQUENCE 444 AA; 49011 MW; 0481ABC4FE5427F8 CRC64;

Query Match 51.3%; Score 101; DB 1; Length 444;
Best Local Similarity 46.3%; Pred. No. 2; Re-09;
Matches 19; Conservative 5; Mismatches 17; Indels 0; Caps 0;

QY 1 ANSFLXLRQSLRXKXLCIXICDFXXKXKIFEDVDDTLAFW 41
Db 40 ANSFLXLRQSLRXKXLCIXICDFXXKXKIFEDVDDTLAFW 80
RESULT 12
P77_HUMAN
ID P77_HUMAN STANDARD; PRT; 466 AA.
AC P08709; Q14339;
DT 01-JAN-1988 (Rel. 06, Created)
DT 01-JAN-1988 (Rel. 06, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Coagulation factor VII precursor (EC 3.4.21.21) (Serum prothrombin
conversion accelerator) (Eptacog alfa).
GN F7.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RX MEDLINE=86205965; PubMed=3486420;
RA Hagen F.S., Gray C.L., O'Hara P.J., Grant F.J., Saari G.C.,
RA Woodbury R.G., Hart C.E., Insley M.Y., Kisiel W., Kurachi K.,
RA Davie E.W.;
RT "Characterization of a cDNA coding for human factor VII.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:2412-2416(1986).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=87260948; PubMed=3037537;
RA O'Hara P.J., Grant F.J., Haldeman B.A., Gray C.L., Insley M.Y.,
RA Hagen F.S., Murray M.J.;
RT "Nucleotide sequence of the gene coding for human factor VII, a
vitamin K-dependent protein participating in blood coagulation.";
RL Proc. Natl. Acad. Sci. U.S.A. 84:5158-5162(1987).
RN [3]
RP SEQUENCE FROM N.A., AND VARIANTS THR-352; GLN-413 AND LYS-445.
RA Rieder M.J., Axel T.Z., Carrington D.P., Chung M.-W., Lee K.L.,
RA Poel C.L., Toth E.J., Yi Q., Nickerson D.A.;
RL Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE OF 61-466, AND POST-TRANSLATIONAL MODIFICATIONS.
RX MEDLINE=89088153; PubMed=3264725;
RA Thim L., Bjorn S., Christensen M., Nicolaissen E.M., Lund-Hansen T.,
RA Pedersen A.H., Hedner U.;
RT "Amino acid sequence and posttranslational modifications of human
factor VIIa from plasma and transfected baby hamster kidney cells.";
RL Biochemistry 27:7785-7793(1988).
RN [5]
RP CARBOHYDRATE-LINKAGE SITES SER-112 AND SER-120.
RX MEDLINE=91250411; PubMed=1904059;
RA Bjorn S., Foster D.C., Thim L., Wiberg F.C., Christensen M.,
RA Koniya Y., Pedersen A.H., Kisiel W.;
RT "Human plasma and recombinant factor VII. Characterization of O-
glycosylations at serine residues 52 and 60 and effects of site-
directed mutagenesis of serine 52 to alanine.";
RL J. Biol. Chem. 266:11051-11057(1991).
RN [6]
RP STRUCTURE OF CARBOHYDRATE ON SER-112.
RX MEDLINE=90062160; PubMed=2511201;
RA Nishimura H., Kawabata S., Kisiel W., Hase S., Ikenaka T., Takao T.,
RA Shimonishi Y., Iwanaga S.;
RT "Identification of a disaccharide (Xyl-Glc) and a trisaccharide
(Xyl2-Glc) O-glycosidically linked to a serine residue in the first
epidermal growth factor-like domain of human factors VII and IX and
protein Z and bovine protein Z.";
RL J. Biol. Chem. 264:20320-20325(1989).
RN [7]
RP STRUCTURE OF CARBOHYDRATE ON SER-112.
RX MEDLINE=91344709; PubMed=2129367;
RA Iwanaga S., Nishimura H., Kawabata S., Kisiel W., Hase S., Ikenaka T.;
RT "A new trisaccharide sugar chain linked to a serine residue in the

RT first BGF-like domain of clotting factors VII and IX and protein Z.";
 RL Adv. Exp. Med. Biol. 281:121-131(1990).
 RN [8]
 RP X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS) OF FVIIA IN COMPLEX WITH TF.
 RX BANNER D.W., D'Arcy A., Chene C., Winkler F.K., Guha A.,
 RA Konigsberg W.H., Nemerson Y., Kirchhofer D.;
 RA "The crystal structure of the complex of blood coagulation factor
 RT VIIa with soluble tissue factor.";
 RL Nature 380:41-46(1996).
 RN [9]
 RP X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF FVIIA IN COMPLEX WITH TF.
 RX MEDLINE=99126538; PubMed=9925787;
 RA Zhang E., St Charles R., Tulinsky A.;
 RA "Structure of extracellular tissue factor complexed with factor VIIa
 RT inhibited with a BPTI mutant.";
 RL J. Mol. Biol. 285:2089-2104(1999).
 RN [10]
 RP STRUCTURE BY NMR OF 105-145.
 RX MEDLINE=98367502; PubMed=9692950;
 RA Muranyi A., Finn B.E., Gippert G.P., Forsen S., Stenflo J.,
 RA Drakenberg T.;
 RA "Solution structure of the N-terminal BGF-like domain from human
 RT factor VII.";
 RL Biochemistry 37:10605-10615(1998).
 RN [11]
 RP VARIANT GLN-364.
 RX MEDLINE=91300046; PubMed=20700047;
 RA O'Brien D.P., Gale K.M., Anderson J.S., McVey J.H., Miller G.J.,
 RA Meade T.W., Tuddenham E.G.D.;
 RA "Purification and characterization of factor VII 304-Gln: a variant
 RT molecule with reduced activity isolated from a clinically unaffected
 RL male.";
 RL Blood 78:132-140(1991).
 RN [12]
 RP VARIANTS GLN-364 AND PHE-370.
 RX MEDLINE=92340074; PubMed=1634227;
 RA Marchetti G., Patrachini P., Gemmati D., Derosa V., Pinotti M.,
 RA Roderigo G., Casonato A., Girolami A., Bernardi F.;
 RA "Detection of two missense mutations and characterization of a repeat
 RT polymorphism in the factor VII gene (F7)." ;
 RL Hum. Genet. 89:497-502(1992).
 RN [13]
 RP VARIANT TYR-238.
 RX MEDLINE=93372811; PubMed=8364544;
 RA Marchetti G., Ferrati M., Patrachini P., Redaelli R., Bernardi F.;
 RA "A missense mutation (178Cys-->Tyr) and two neutral dimorphisms
 RT (115His and 333Ser) in the human coagulation factor VII gene." ;
 RL Hum. Mol. Genet. 2:1055-1056(1993).
 RN [14]
 RP VARIANTS.
 RX MEDLINE=94061028; PubMed=8242057;
 RA Takamiya O., Kemball-Cook G., Martin D.M.A., Cooper D.N.,
 RA von Felten A., Medli E., Hahn I., Pragnell D.R., Lumley H.,
 RA Tuddenham E.G.D., McVey J.H.;
 RA "Detection of missense mutations by single-strand conformational
 RT polymorphism (SSCP) analysis in five dysfunctional variants of
 RL coagulation factor VII." ;
 RL Hum. Mol. Genet. 2:1355-1359(1993).
 RN [15]
 RP VARIANTS CHARLOTTE GLN-139 AND GLN-212.
 RX MEDLINE=94264305; PubMed=8204879;
 RA Chaing S., Clarke B., Sridhara S., Chu K., Friedman P., Vandusen W.,
 RA Roberts H.R., Blajchman M., Monroe D.M., High K.A.;
 RA "Severe factor VII deficiency caused by mutations abolishing the
 RT cleavage site for activation and altering binding to tissue factor." ;
 RL Blood 83:3524-3535(1994).
 RN [16]
 RP VARIANT VAL-354.
 RX MEDLINE=95072589; PubMed=7981691;
 RA Bernardi F., Castaman G., Redaelli R., Pinotti M., Lunghi B.,
 RA Rodeghiero F., Marchetti G.;
 RA "Topologically equivalent mutations causing dysfunctional coagulation

RT factors VII (294Ala-->Val) and X (334Ser-->Pro)." ;
 RL Hum. Mol. Genet. 3:1175-1177(1994).
 RN [17]
 RP VARIANT MIE HIS-307.
 RX MEDLINE=95064662; PubMed=7974346;
 RA Ohiwa M., Hayashi T., Wada H., Minamikawa K., Shirakawa S.,
 RA Suzuki K.;
 RA "Factor VII Mie: homozygous asymptomatic type I deficiency caused by
 RT an amino acid substitution of His (CAC) for Arg(247) (CGC) in the
 RL catalytic domain." ;
 RL Thromb. Haemost. 71:773-777(1994).
 RN [18]
 RP VARIANT MET-419.
 RX MEDLINE=96247510; PubMed=8652821;
 RA Arbini A.A., Mannucci P.M., Bader K.A.;
 RA "A Thr39Met mutation in factor VII of a patient with a hereditary
 RT deficiency causes defective secretion of the molecule." ;
 RL Blood 87:5085-5094(1996).
 RN [19]
 RP VARIANTS TRP-283; LYS-325; VAL-358; GLN-364; GLU-402 AND GLN-413.
 RX MEDLINE=97001216; PubMed=8944208;
 RA Bernardi F., Castaman G., Pinotti M., Ferraresi P., di Iasio M.G.,
 RA Lunghi B., Rodeghiero F., Marchetti G.;
 RA "Mutation pattern in clinically asymptomatic coagulation factor VII
 RT deficiency." ;
 RL Hum. Mutat. 8:108-115(1996).
 RN [20]
 RP VARIANT VAL-304.
 RX MEDLINE=97037613; PubMed=8883260;
 RA Tamary H., Fromovich Y., Shalom D., Reich Z., Dym O., Lanir N.,
 RA Brenner B., Paz M., Luder A.S., Blau O., Korostishevsky M.,
 RA Zaitov R., Seligsohn U.;
 RA "Ala244Val is a common, probably ancient mutation causing factor VII
 RT deficiency in Moroccan and Iranian Jews." ;
 RL Thromb. Haemost. 76:283-291(1996).
 RN [21]
 RP VARIANT MORIOKA PRO-13.
 RX MEDLINE=98235713; PubMed=9576180;
 RA Ozawa T., Takikawa Y., Niliya K., Ejiri N., Suzuki K., Sato S.,
 RA Sakuragawa N.;
 RA "Factor VII Moriooka (FVII L-26P): a homozygous missense mutation in
 RT the signal sequence identified in a patient with factor VII
 RL deficiency." ;
 RL Br. J. Haematol. 101:47-49(1998).
 RN [22]
 RP VARIANTS MALTA THR-194 AND VAL-304.
 RX MEDLINE=98112461; PubMed=9452082;
 RA Alshinawi C., Scerri C., Galdies R., Aquilina A., Felice A.E.;
 RA "Two new missense mutations (P134T and A244V) in the coagulation
 RT factor VII gene." ;
 RL Hum. Mutat. Suppl. 1:S189-S191(1998).
 RN [23]
 RP VARIANTS ASP-295 AND GLN-413.
 RX MEDLINE=99318093; PubMed=10391209;
 RA Cagill M., Altschuler D., Ireland J., Sklar P., Ardlie K., Patil N.,
 RA Shaw N., Lane C.R., Lim E.P., Kalyanaram N., Nimesh J., Ziaugra L.,
 RA Friedland L., Rolfe A., Warrington J., Lipschutz R., Daley G.Q.,
 RA Lander E.S.;
 RA "Characterization of single-nucleotide polymorphisms in coding regions
 RT of human genes." ;
 RL Nat. Genet. 22:231-238(1999).
 RN [24]
 RP Query Match 50.3%; Score 99; DB 1; Length 466;
 RL Best Local Similarity 48.8%; Pred. No. 6:5e-09;
 RN Matches 20; Conservative 4; Mismatches 17; Indels 0; Gaps 0;
 RN QY 1 ANSPFLXLRGSLRXRCIXXICDFXXKXIFEDVDDTLAFW 41
 RL 61 ANAFLELRPGSLERECKEQCSFEAREFKDAERTKLEW 101

RESULT 13
 TMGL_HUMAN


```

DR SMART; SM00130; KR; 2.
DR SMART; SM00020; TRYP_SPC; 1.
DR PROSITE; PS00011; GLU CARBOXYLATION; 1.
DR PROSITE; PS00021; KRINGLE_1; 2.
DR PROSITE; PS00070; KRINGLE_2; 2.
DR PROSITE; PS00240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00135; TRYPSIN_SER; 1.
KW Blood coagulation; Plasma; Calcium-binding; Glycoprotein; Repeat;
KW Vitamin K; Zymogen; Gamma-carboxylglutamic acid; Acute phase; Liver;
KW Hydrolase; Serine protease; Kringle; Signal.
FT SIGNAL 1 24
FT PROPEP 25 43
FT CHAIN 44 617
FT PEPTIDE 44 200
FT PEPTIDE 201 323
FT CHAIN 324 359
FT CHAIN 360 617
FT DOMAIN 109 187
FT DOMAIN 215 292
FT DOMAIN 360 617
FT SITE 200 201
FT SITE 323 324
FT SITE 359 360
FT ACT_SITE 402 402
FT ACT_SITE 458 458
FT ACT_SITE 564 564
FT MOD_RES 50 50
FT MOD_RES 51 51
FT MOD_RES 58 58
FT MOD_RES 60 60
FT MOD_RES 63 63
FT MOD_RES 64 64
FT MOD_RES 69 69
FT MOD_RES 70 70
FT MOD_RES 73 73
FT MOD_RES 76 76
FT CARBOHYD 120 120
FT CARBOHYD 144 144
FT CARBOHYD 412 412
FT CARBOHYD 552 552
FT DISULFID 61 66
FT DISULFID 91 104
FT DISULFID 109 187
FT DISULFID 130 170
FT DISULFID 158 182
FT DISULFID 215 292
FT DISULFID 236 276
FT DISULFID 264 287
FT DISULFID 332 478
FT DISULFID 387 403
FT DISULFID 532 546
FT DISULFID 560 590
SQ SEQUENCE 617 AA; AD27DLB17445DB1D CRC64;

Query Match 43.9%; Score 86.5; DB 1; Length 617;
Best Local Similarity 42.2%; Pred. No. 1.2e-06;
Matches 19; Conservative 6; Mismatches 19; Indels 1; Gaps 1;

OY 1 ANS-FLKXLFQGSIXEXXCIXXICDFXXKXIFEDVDDTLAFWSKH 44
Db 44 ANSGFLEELRKNGLRECEVEEQCSYEAFEALESPOQTDVFWAKY 88

RESULT 15
THRB_MOUSE
ID THRB_MOUSE STANDARD; PRT; 618 AA.
AC P19221;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE Prothrombin precursor (EC 3.4.21.5).
GN F2 OR CF2.

```

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OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NCBI_TaxID=10090;
RN [1]
RN SEQUENCE FROM N.A.
RP STRAIN=C57BL/6; TISSUE=Liver;
RC MEDLINE=91025551; PubMed=222810;
RX Friesner Degen S.J., Schaffer L.A., Jamison C.S., Grant S.G.,
RA Fitzgibbon J.J., Fai J.A., Chapman V.M., Elliott R.W.;
RT "Characterization of the cDNA coding for mouse prothrombin and
RT localization of the gene on mouse chromosome 2.";
RL DNA Cell Biol. 9:487-498(1990).
RN [2]
RN SEQUENCE FROM N.A.
RP STRAIN=FVB/N; TISSUE=Liver;
RC MEDLINE=22388257; PubMed=12477932;
RX Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickinson M.C.,
RA Rodriguez A.C., Greenwood J., Schmutz J., Myers R.M.,
RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smalls D.E.,
RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [3]
RN SEQUENCE OF 384-618 FROM N.A.
RP TISSUE=Liver;
RC MEDLINE=92212913; PubMed=1557383;
RX Banfield D.K., Macgillivray R.T.;
RT "Partial characterization of vertebrate prothrombin cDNAs:
RT amplification and sequence analysis of the B chain of thrombin from
RT nine different species.";
RL Proc. Natl. Acad. Sci. U.S.A. 89:2779-2783(1992).
CC -!- FUNCTION: Thrombin, which cleaves bonds after Arg and Lys,
CC converts fibrinogen to fibrin and activates factors V, VII, VIII,
CC XIII, and, in complex with thrombomodulin, protein C.
CC -!- CATALYTIC ACTIVITY: Preferential cleavage: Arg-|-Gly; activates
CC fibrinogen to fibrin and releases fibrinopeptide A and B.
CC -!- PTM: THE GAMMA-CARBOXYGLUTAMYL RESIDUES, WHICH BIND CALCIUM IONS,
CC RESULT FROM THE CARBOXYLATION OF GLUTAMYL RESIDUES BY A MICROSOMAL
CC ENZYME, THE VITAMIN K-DEPENDENT CARBOXYLASE. THE MODIFIED RESIDUES
CC ARE NECESSARY FOR THE CA-DEPENDENT INTERACTION WITH A NEGATIVELY
CC CHARGED PHOSPHOLIPID SURFACE, WHICH IS ESSENTIAL FOR THE CONVERSION
CC OF PROTHROMBIN TO THROMBIN.
CC -!- MISCELLANEOUS: Prothrombin is activated on the surface of a
CC phospholipid membrane that binds the amino end of prothrombin and
CC factors Va and Xa in Ca-dependent interactions; factor Xa removes
CC the activation peptide and cleaves the remaining part into light
CC and heavy chains. The activation process starts slowly because
CC factor V itself has to be activated by the initial, small amounts
CC of thrombin.
CC -!- MISCELLANEOUS: Thrombin can itself cleave the amino terminal
CC fragment (fragment 1) of the prothrombin, prior to its activation
CC by factor Xa.
CC -!- SIMILARITY: Belongs to peptidase family S1.
CC -!- SIMILARITY: Contains 2 kringle domains.
CC -----
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CC the European Bioinformatics Institute. There are no restrictions on its

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CC EMBL; X52308; CAA36548.1; .
CC EMBL; BC013662; AAH13662.1; .
CC EMBL; M81394; AAA40435.1; .
CC PIR; A35827; A35827.
CC HSP; P00734; 1B7X.
CC MEROPS; S01.217; .
CC MGD; MGI:88180; F2.
CC InterPro; IPR009003; Cys Ser trypsin.
CC InterPro; IPR002383; GLA blood.
CC InterPro; IPR000001; Kringle.
CC InterPro; IPR001254; Peptidase S1.
CC InterPro; IPR001314; Peptidase S1A.
CC InterPro; IPR003966; Peptidase S1A pr.
CC InterPro; IPR000294; VitK_dep_GLA.
CC Pfam; PF00594; Gla; 1.
CC Pfam; PF00051; kringle; 2.
CC Pfam; PF00089; trypsin; 1.
CC PRINTS; PR00722; CHYMOTRYPSIN.
CC PRINTS; PR00001; GLABLOOD.
CC PRINTS; PR00018; KRINGLE.
CC PRINTS; PR01505; PROTHROMBIN.
CC ProDom; PD000395; Kringle; 2.
CC SMART; SM00069; GLA; 1.
CC SMART; SM00130; KR; 2.
CC SMART; SM00020; Tryp SPC; 1.
CC PROSITE; PS00011; GLU CARBOXYLATION; 1.
CC PROSITE; PS00021; KRINGLE 1; 2.
CC PROSITE; PS00070; KRINGLE 2; 2.
CC PROSITE; PS00240; TRYPSIN_DOM; 1.
CC PROSITE; PS00134; TRYPSIN_HIS; 1.
CC PROSITE; PS00135; TRYPSIN_SER; 1.
KW Blood coagulation; Plasma; Calcium-binding; Glycoprotein; Repeat;
KW Vitamin K; Zymogen; Gamma-carboxyglutamic acid; Acute phase; Liver;
KW Hydroxylase; Serine protease; Kringle; Signal.
FT SIGNAL 1 24
FT PROPEP 25 43
FT CHAIN 44 618
FT PEPTIDE 44 200
FT PEPTIDE 201 324
FT CHAIN 325 360
FT CHAIN 361 618
FT DOMAIN 109 187
FT DOMAIN 215 292
FT DOMAIN 351 618
FT SITE 200 201
FT SITE 324 325
FT SITE 360 361
FT ACT_SITE 403 403
FT ACT_SITE 459 459
FT ACT_SITE 565 565
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FT MOD_RES 73 73
FT MOD_RES 76 76
FT DISULFID 61 66
FT DISULFID 91 104
FT DISULFID 109 187
FT DISULFID 130 170
FT DISULFID 158 182
FT DISULFID 215 293
FT DISULFID 236 276
FT DISULFID 264 288

FT DISULFID 333 479 INTERCHAIN (BY SIMILARITY).
FT DISULFID 388 404 BY SIMILARITY.
FT DISULFID 533 547 BY SIMILARITY.
FT DISULFID 561 591 BY SIMILARITY.
FT CARBOHYD 122 122 N-LINKED (GLCNAC. . .).
FT CARBOHYD 144 144 N-LINKED (GLCNAC. . .).
FT CARBOHYD 413 413 N-LINKED (GLCNAC. . .).
FT CARBOHYD 553 553 N-LINKED (GLCNAC. . .).
SQ SEQUENCE 618 AA; 70268 MW; B89P719AAF601E0 CRC64;
Query Match 43.98; Score 86.5; DB 1; Length 618;
Best Local Similarity 42.28; Pred. No. 1.2e-06;
Matches 19; Conservative 6; Mismatches 19; Indels 1; Gaps 1;
QY 1 ANS-FLXXLRQGSIXXCIXXICDFXAXKXIFEDVDDTLAFWSKH 44
Db 44 ANSGFLEELRKGNLRECVEEQCSYEAFEALESPOQTDVFWAKY 88
Search completed: March 1, 2004, 10:03:23
Job time : 10.5 secs